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- ☐ CONTINUATION-IN-PART
- ☒ CONTINUATION
- ☐ DIVISIONAL
- ☐ FILE WRAPPER CONTINUATION

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Sir:

Transmitted herewith for filing is the continuation patent application of

Inventors: **Patrick V. Warren and Ronald V. Swanson**

For: **TRANSAMINASES AND AMINOTRANSFERASES**

This is a request for filing a X continuation divisional application under 37 C.F.R. 1.53(b) , of prior application Serial No. 08/646,590, filed on May 8, 1996, now pending, which is entitled **TRANSAMINASES AND AMINOTRANSFERASES**, which is a continuation-in-part of copending U.S. application Serial No. 08/599,171 filed February 9, 1996.

by the following named inventors: Patrick V. Warren and Ronald V. Swanson

FULL NAME OF FIRST INVENTOR	LAST NAME: Warren	FIRST NAME: Patrick	MIDDLE NAME: V.
CITIZENSHIP	STATE OR FOREIGN COUNTRY: United States		
POST OFFICE ADDRESS	POST OFFICE ADDRESS: 3507 Sheffield Avenue	CITY AND STATE: Philadelphia, Pennsylvania	ZIP CODE: 19163
FULL NAME OF SECOND INVENTOR	LAST NAME: Swanson	FIRST NAME: Ronald	MIDDLE NAME: V.
CITIZENSHIP	STATE OR FOREIGN COUNTRY: United States		
POST OFFICE ADDRESS:	POST OFFICE ADDRESS: 309 No. Lemon Street, Apt. A	CITY AND STATE: Media, Pennsylvania	ZIP CODE: 19063

No payment of the issue fee, abandonment of, or termination of proceeding has occurred in the above-identified prior application.

9. Please transfer the drawings from the prior application to the new application.
10. X A true copy of the prior application as filed is enclosed, including the Declaration and Power of Attorney filed in parent application, U.S. Serial No. 08/599,171 filed February 9, 1996, the Substitute Declaration and Power of Attorney filed December 26, 1996, and the Revocation and New Power of Attorney filed on June 13, 1997.
11. X No Information Disclosure Statement(s) was filed in the prior application under 37 C.F.R. 1.97.
12. Also enclosed: Copy of Petition for Extension of Time in parent application U.S. Serial No.:

Address all future communications to:

Lisa A. Haile, Ph.D.
GRAY CARY WARE & FREIDENRICH LLP
4365 Executive Drive, Suite 1600
San Diego, California 92121-2189
Telephone: 619-677-1456
Facsimile: 619-677-1465

The undersigned states that the enclosed application papers comprise a true copy of the prior application as filed.

Respectfully submitted,

Date: 9/2/99



Lisa A. Haile, Ph.D.
Attorney for Applicant
Registration No. 38,347

GRAY CARY WARE & FREIDENRICH LLP
4365 Executive Drive, Suite 1600
San Diego, CA 92121-2189

TRANSAMINASES AND AMINOTRANSFERASES

This application is a continuation-in-part of copending U.S. serial no. 08/599,171 filed on February 9, 1996.

This invention relates to newly identified polynucleotides, polypeptides encoded by such polynucleotides, the use of such polynucleotides and polypeptides, as well as the production and isolation of such polynucleotides and polypeptides. More particularly, the polynucleotides and polypeptides of the present invention have been putatively identified as transaminases and/or aminotransferases. Aminotransferases are enzymes that catalyze the transfer of amino groups from α -amino to α -keto acids. They are also called transaminases.

The α -amino groups of the 20 L-amino acids commonly found in proteins are removed during the oxidative degradation of the amino acids. The removal of the α -amino groups, the first step in the catabolism of most of the L-amino acids, is promoted by aminotransferases (or transaminases). In these transamination reactions, the α -amino group is transferred to the α -carbon atom of α -ketoglutarate, leaving behind the corresponding α -keto acid analog of the amino acid. There is no net deamination (*i.e.*,

loss of amino groups) in such reactions because the α -ketoglutarate becomes aminated as the α -amino acid is deaminated. The effect of transamination reactions is to collect the amino groups from many different amino acids in the form of only one, namely, L-glutamate. The glutamate channels amino groups either into biosynthetic pathways or into a final sequence of reactions by which nitrogenous waste products are formed and then excreted.

Cells contain several different aminotransferases, many specific for α -ketoglutarate as the amino group acceptor. The aminotransferases differ in their specificity for the other substrate, the L-amino acid that donates the amino group, and are named for the amino group donor. The reactions catalyzed by the aminotransferases are freely reversible, having an equilibrium constant of about 1.0 ($\Delta G^0 \approx 0$ kJ/mol).

Aminotransferases are classic examples of enzymes catalyzing bimolecular ping-pong reactions. In such reactions the first substrate must leave the active site before the second substrate can bind. Thus the incoming amino acid binds to the active site, donates its amino group to pyridoxal phosphate, and departs in the form of an α -keto acid. Then the incoming α -keto acid is bound, accepts the amino group from pyridoxamine phosphate, and departs in the form of an amino acid.

The measurement of alanine aminotransferase and aspartate aminotransferase levels in blood serum is an important diagnostic procedure in medicine, used as an indicator of heart damage and to monitor recovery from the damage.

The polynucleotides and polypeptides of the present invention have been identified as transaminases and/or aminotransferases as a result of their enzymatic activity.

In accordance with one aspect of the present invention, there are provided novel enzymes, as well as active fragments, analogs and derivatives thereof.

In accordance with another aspect of the present invention, there are provided isolated nucleic acid molecules encoding the enzymes of the present invention including mRNAs, cDNAs, genomic DNAs as well as active analogs and fragments of such enzymes.

In accordance with another aspect of the present invention there are provided isolated nucleic acid molecules encoding mature polypeptides expressed by the DNA contained in ATCC Deposit No. _____.

In accordance with yet a further aspect of the present invention, there is provided a process for producing such polypeptides by recombinant techniques comprising culturing recombinant prokaryotic and/or eukaryotic host cells, containing a nucleic acid sequence of the present invention, under conditions promoting expression of said enzymes and subsequent recovery of said enzymes.

In accordance with yet a further aspect of the present invention, there is provided a process for utilizing such enzymes, or polynucleotides encoding such enzymes for transferring an amino group from an α -amino acid to an α -keto acid. Most transaminases use L-amino acids as substrates, but as described below, it is also possible to convert the transaminases of the invention to use D-amino acids as substrates, thereby increasing their array of uses to include, for example, manufacture of synthetic pyrethroids and as components of β -lactam antibiotics. The transaminases of the invention are stable at high temperatures and in organic solvents and, thus, are superior for use with L- and/or D-amino acids for production of optically pure chiral compounds used in pharmaceutical, agricultural and other chemical industries.

In accordance with yet a further aspect of the present invention, there are also provided nucleic acid probes comprising nucleic acid molecules of sufficient length to hybridize to a nucleic acid sequence of the present invention.

In accordance with yet a further aspect of the present invention, there is provided a process for utilizing such enzymes, or polynucleotides encoding such enzymes, for *in vitro* purposes related to scientific research, for example, to generate probes for identifying similar sequences which might encode similar enzymes from other organisms by using certain regions, *i.e.*, conserved sequence regions, of the nucleotide sequence.

These and other aspects of the present invention should be apparent to those skilled in the art from the teachings herein.

The following drawings are illustrative of embodiments of the invention and are not meant to limit the scope of the invention as encompassed by the claims.

Figure 1 is an illustration of the full-length DNA (SEQ ID NO:17) and corresponding deduced amino acid sequence (SEQ ID NO:25) of *Aquifex* aspartate transaminase A of the present invention. Sequencing was performed using a 378 automated DNA sequencer (Applied Biosystems, Inc.) for all sequences of the present invention.

Figure 2 is an illustration of the full-length DNA (SEQ ID NO:18) and corresponding deduced amino acid sequence (SEQ ID NO:26) of *Aquifex* aspartate aminotransferase B.

Figure 3 is an illustration of the full-length DNA (SEQ ID NO:19) and corresponding deduced amino acid sequence (SEQ ID NO:27) of *Aquifex* adenosyl-8-amino-7-oxononanoate aminotransferase.

Figure 4 is an illustration of the full-length DNA (SEQ ID NO:20) and corresponding deduced amino acid sequence (SEQ ID NO:28) of *Aquifex* acetylornithine aminotransferase.

Figure 5 is an illustration of the full-length DNA (SEQ ID NO:21) and corresponding deduced amino acid sequence (SEQ ID NO:29) of *Ammonifex degensii* aspartate aminotransferase.

Figure 6 is an illustration of the full-length DNA (SEQ ID NO:22) and corresponding deduced amino acid sequence (SEQ ID NO:30) of *Aquifex* glucosamine:fructose-6-phosphate aminotransferase.

Figure 7 is an illustration of the full-length DNA (SEQ ID NO:23) and corresponding deduced amino acid sequence (SEQ ID NO:31) of *Aquifex* histidinol-phosphate aminotransferase.

Figure 8 is an illustration of the full-length DNA (SEQ ID NO:24) and corresponding deduced amino acid sequence (SEQ ID NO:32) of *Pyrobaculum aerophilum* branched chain aminotransferase.

Figure 9 is a diagrammatic illustration of the assay used to assess aminotransferase activity of the proteins using glutamate dehydrogenase.

The term "gene" means the segment of DNA involved in producing a polypeptide chain; it includes regions preceding and following the coding region (leader and trailer) as well as intervening sequences (introns) between individual coding segments (exons).

A coding sequence is "operably linked to" another coding sequence when RNA polymerase will transcribe the two coding sequences into a single mRNA, which is then translated into a single polypeptide having amino acids derived from both coding sequences. The coding sequences need not be contiguous to one another so long as the expressed sequences ultimately process to produce the desired protein.

"Recombinant" enzymes refer to enzymes produced by recombinant DNA techniques; *i.e.*, produced from cells transformed by an exogenous DNA construct encoding the desired enzyme. "Synthetic" enzymes are those prepared by chemical synthesis.

A DNA "coding sequence of" or a "nucleotide sequence encoding" a particular enzyme, is a DNA sequence which is transcribed and translated into an enzyme when placed under the control of appropriate regulatory sequences.

In accordance with an aspect of the present invention, there are provided isolated nucleic acids (polynucleotides) which encode for the mature enzymes having the deduced amino acid sequences of Figures 1-8 (SEQ ID NOS:17-32).

In accordance with another aspect of the present invention, there are provided isolated polynucleotides encoding the enzymes of the present invention. The deposited material is a mixture of genomic clones comprising DNA encoding an enzyme of the present invention. Each genomic clone comprising the respective DNA has been inserted into a pQE vector (Quiagen, Inc., Chatsworth, CA). The deposit has been deposited with the American Type Culture Collection, 12301 Parklawn Drive, Rockville, Maryland 20852, USA, on December 13, 1995 and assigned ATCC Deposit No. _____.

The deposit(s) have been made under the terms of the Budapest Treaty on the International Recognition of the deposit of micro-organisms for purposes of patent procedure. The strains will be irrevocably and without restriction or condition released to the public upon the issuance of a patent. These deposits are provided merely as convenience to those of skill in the art and are not an admission that a deposit would be required under 35 U.S.C. §112. The sequences of the polynucleotides contained in the deposited materials, as well as the amino acid sequences of the polypeptides encoded thereby, are controlling in the event of any conflict with any description of sequences

herein. A license may be required to make, use or sell the deposited materials, and no such license is hereby granted.

The polynucleotides of this invention were originally recovered from genomic DNA libraries derived from the following organisms:

Aquifex VF5 is a Eubacteria which was isolated in Vulcano, Italy. It is a gram-negative, rod-shaped, strictly chemolithoautotrophic, marine organism which grows optimally at 85-90°C ($T_{max}=95^{\circ}\text{C}$) at pH 6.8 in a high salt culture medium with O_2 as a substrate, and $\text{H}_2/\text{CO}_2+0.5\%$ O_2 in gas phase.

Ammonifex degensii KC4 is a new Eubacaterial organism isolated in Java, Indonesia. This Gram negative chemolithoautotroph has three respiration systems. The bacterium can utilize nitrate, sulfate, and sulfur. The organism grows optimally at 70°C, and pH 7.0, in a low salt culture medium with 0.2% nitrate as a substrate and H_2/CO_2 in gas phase.

Pyrobaculum aerophilum IM2 is a thermophilic sulfur archaea (Crenarchaeota) isolated in Ischia Maronti, Italy. It is a rod-shaped organism that grows optimally at 100°C at pH 7.0 in a low salt culture medium with nitrate, yeast extract, peptone, and O_2 as substrates and N_2/CO_2 , O_2 in gas phase.

Accordingly, the polynucleotides and enzymes encoded thereby are identified by the organism from which they were isolated, and are sometimes hereinafter referred to as "VF5/ATA" (Figure 1 and SEQ ID NOS:17 and 25), "VF5/AAB" (Figure 2 and SEQ ID NOS:18 and 26), "VF5/A87A" (Figure 3 and SEQ ID NOS:19 and 27), "VF5/AOA" (Figure 4 and SEQ ID NOS:20 and 28), "KC4/AA" (Figure 5 and SEQ ID NOS:21 and 29), "VF5/GF6PA" (Figure 6 and SEQ ID NOS:22 and 30), "VF5/HPA" (Figure 7 and SEQ ID NOS:23 and 31) and "IM2/BCA" (Figure 8 and SEQ ID NOS:24 and 32).

The polynucleotides and polypeptides of the present invention show identity at the nucleotide and protein level to known genes and proteins encoded thereby as shown in Table 1.

Table 1

Enzyme	Gene w/closest Homology (Organism)	Protein Similarity (%)	Protein Identity (%)	DNA Identity (%)
VF5/ATA	<i>Bacillus subtilis</i>	57.5	38.3	50.1
VF5/AAB	<i>Sulfolobus solfataricus</i>	62.5	33.0	50.1
VF5/A87A	<i>Bacillus sphaericus BioA</i>	67.4	42.9	51
VF5/AOA	<i>Bacillus subtilis argD</i>	70.6	48.7	52.0
KC4/AA	<i>Bacillus YM-2 aspC</i>	72.6	52.7	52.0
VF5/GF6PA	<i>Rhizobium Leguminosarum NodM</i>	66.3	47.7	51.0
VF5/HPA	<i>Bacillus subtilis HisH/E.coli HisC (same gene)</i>	55.7	32.6	45.3
IM2/BCA	<i>E.coli iluE</i>	63.7	43.6	49.7

All the clones identified in Table 1 encode polypeptides which have transaminase or aminotransferase activity.

One means for isolating the nucleic acid molecules encoding the enzymes of the present invention is to probe a gene library with a natural or artificially designed probe using art recognized procedures (see, for example: Current Protocols in Molecular Biology, Ausubel F.M. *et al.* (EDS.) Green Publishing Company Assoc. and John Wiley Interscience, New York, 1989, 1992). It is appreciated by one skilled in the art that the polynucleotides of SEQ ID NOS:17-24, or fragments thereof (comprising at least 12 contiguous nucleotides), are particularly useful probes. Other particularly

useful probes for this purpose are hybridizable fragments of the sequences of SEQ ID NOS:1-9 (*i.e.*, comprising at least 12 contiguous nucleotides).

With respect to nucleic acid sequences which hybridize to specific nucleic acid sequences disclosed herein, hybridization may be carried out under conditions of reduced stringency, medium stringency or even stringent conditions. As an example of oligonucleotide hybridization, a polymer membrane containing immobilized denatured nucleic acids is first prehybridized for 30 minutes at 45°C in a solution consisting of 0.9 M NaCl, 50 mM NaH₂PO₄, pH 7.0, 5.0 mM Na₂EDTA, 0.5% SDS, 10X Denhardt's, and 0.5 mg/mL polyriboadenylic acid. Approximately 2 X 10⁷ cpm (specific activity 4-9 X 10⁸ cpm/ug) of ³²P end-labeled oligonucleotide probe are then added to the solution. After 12-16 hours of incubation, the membrane is washed for 30 minutes at room temperature in 1X SET (150 mM NaCl, 20 mM Tris hydrochloride, pH 7.8, 1 mM Na₂EDTA) containing 0.5% SDS, followed by a 30 minute wash in fresh 1X SET at T_m -10°C (T_m is minus 10°C) for the oligo-nucleotide probe. The membrane is then exposed to auto-radiographic film for detection of hybridization signals.

Stringent conditions means hybridization will occur only if there is at least 90% identity, preferably at least 95% identity and most preferably at least 97% identity between the sequences. See J. Sambrook *et al.*, *Molecular Cloning, A Laboratory Manual, 2d Ed.*, Cold Spring Harbor Laboratory (1989) which is hereby incorporated by reference in its entirety.

As used herein, a first DNA (RNA) sequence is at least 70% and preferably at least 80% identical to another DNA (RNA) sequence if there is at least 70% and preferably at least a 80% or 90% identity, respectively, between the bases of the first sequence and the bases of the another sequence, when properly aligned with each other, for example when aligned by BLASTN.

The present invention relates to polynucleotides which differ from the reference polynucleotide such that the changes are silent changes, for example the change does not or the changes do not alter the amino acid sequence encoded by the polynucleotide. The present invention also relates to nucleotide changes which result in amino acid substitutions, additions, deletions, fusions and truncations in the polypeptide encoded by the reference polynucleotide. In a preferred aspect of the invention these polypeptides retain the same biological action as the polypeptide encoded by the reference polynucleotide.

The polynucleotides of this invention were recovered from genomic gene libraries from the organisms listed in Table 1. Gene libraries were generated in the Lambda ZAP II cloning vector (Stratagene Cloning Systems). Mass excisions were performed on these libraries to generate libraries in the pBluescript phagemid. Libraries were generated and excisions were performed according to the protocols/methods hereinafter described.

The polynucleotides of the present invention may be in the form of RNA or DNA which DNA includes cDNA, genomic DNA, and synthetic DNA. The DNA may be double-stranded or single-stranded, and if single stranded may be the coding strand or non-coding (anti-sense) strand. The coding sequences which encodes the mature enzymes may be identical to the coding sequences shown in Figures 1-8 (SEQ ID NOS:17-24) or may be a different coding sequence which coding sequence, as a result of the redundancy or degeneracy of the genetic code, encodes the same mature enzymes as the DNA of Figures 1-8 (SEQ ID NOS:17-24).

The polynucleotide which encodes for the mature enzyme of Figures 1-8 (SEQ ID NOS:25-32) may include, but is not limited to: only the coding sequence for the mature enzyme; the coding sequence for the mature enzyme and additional coding sequence such as a leader sequence or a proprotein sequence; the coding sequence for the mature enzyme (and optionally additional coding sequence) and non-coding

sequence, such as introns or non-coding sequence 5' and/or 3' of the coding sequence for the mature enzyme.

Thus, the term "polynucleotide encoding an enzyme (protein)" encompasses a polynucleotide which includes only coding sequence for the enzyme as well as a polynucleotide which includes additional coding and/or non-coding sequence.

The present invention further relates to variants of the hereinabove described polynucleotides which encode for fragments, analogs and derivatives of the enzymes having the deduced amino acid sequences of Figures 1-8 (SEQ ID NOS:25-32). The variant of the polynucleotide may be a naturally occurring allelic variant of the polynucleotide or a non-naturally occurring variant of the polynucleotide.

Thus, the present invention includes polynucleotides encoding the same mature enzymes as shown in Figures 1-8 (SEQ ID NOS:17-24) as well as variants of such polynucleotides which variants encode for a fragment, derivative or analog of the enzymes of Figures 1-8 (SEQ ID NOS:17-24). Such nucleotide variants include deletion variants, substitution variants and addition or insertion variants.

As hereinabove indicated, the polynucleotides may have a coding sequence which is a naturally occurring allelic variant of the coding sequences shown in Figures 1-8 (SEQ ID NOS:17-24). As known in the art, an allelic variant is an alternate form of a polynucleotide sequence which may have a substitution, deletion or addition of one or more nucleotides, which does not substantially alter the function of the encoded enzyme. Also, using directed and other evolution strategies, one may make very minor changes in DNA sequence which can result in major changes in function.

Fragments of the full length gene of the present invention may be used as hybridization probes for a cDNA or a genomic library to isolate the full length DNA and to isolate other DNAs which have a high sequence similarity to the gene or similar

biological activity. Probes of this type preferably have at least 10, preferably at least 15, and even more preferably at least 30 bases and may contain, for example, at least 50 or more bases. The probe may also be used to identify a DNA clone corresponding to a full length transcript and a genomic clone or clones that contain the complete gene including regulatory and promotor regions, exons and introns. An example of a screen comprises isolating the coding region of the gene by using the known DNA sequence to synthesize an oligonucleotide probe. Labeled oligonucleotides having a sequence complementary or identical to that of the gene or portion of the gene sequences of the present invention are used to screen a library of genomic DNA to determine which members of the library the probe hybridizes to.

It is also appreciated that such probes can be and are preferably labeled with an analytically detectable reagent to facilitate identification of the probe. Useful reagents include but are not limited to radioactivity, fluorescent dyes or enzymes capable of catalyzing the formation of a detectable product. The probes are thus useful to isolate complementary copies of DNA from other sources or to screen such sources for related sequences.

The present invention further relates to polynucleotides which hybridize to the hereinabove-described sequences if there is at least 70%, preferably at least 90%, and more preferably at least 95% identity between the sequences. The present invention particularly relates to polynucleotides which hybridize under stringent conditions to the hereinabove-described polynucleotides. As herein used, the term "stringent conditions" means hybridization will occur only if there is at least 95% and preferably at least 97% identity between the sequences. The polynucleotides which hybridize to the hereinabove described polynucleotides in a preferred embodiment encode enzymes which either retain substantially the same biological function or activity as the mature enzyme encoded by the DNA of Figures 1-8 (SEQ ID NOS:17-24).

Alternatively, the polynucleotide may have at least 15 bases, preferably at least 30 bases, and more preferably at least 50 bases which hybridize to any part of a polynucleotide of the present invention and which has an identity thereto, as hereinabove described, and which may or may not retain activity. For example, such polynucleotides may be employed as probes for the polynucleotides of SEQ ID NOS:17-24, for example, for recovery of the polynucleotide or as a diagnostic probe or as a PCR primer.

Thus, the present invention is directed to polynucleotides having at least a 70% identity, preferably at least 90% identity and more preferably at least a 95% identity to a polynucleotide which encodes the enzymes of SEQ ID NOS:25-32 as well as fragments thereof, which fragments have at least 15 bases, preferably at least 30 bases and most preferably at least 50 bases, which fragments are at least 90% identical, preferably at least 95% identical and most preferably at least 97% identical under stringent conditions to any portion of a polynucleotide of the present invention.

The present invention further relates to enzymes which have the deduced amino acid sequences of Figures 1-8 (SEQ ID NOS:17-24) as well as fragments, analogs and derivatives of such enzyme.

The terms "fragment," "derivative" and "analog" when referring to the enzymes of Figures 1-8 (SEQ ID NOS:25-32) means enzymes which retain essentially the same biological function or activity as such enzymes. Thus, an analog includes a proprotein which can be activated by cleavage of the proprotein portion to produce an active mature enzyme.

The enzymes of the present invention may be a recombinant enzyme, a natural enzyme or a synthetic enzyme, preferably a recombinant enzyme.

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The fragment, derivative or analog of the enzymes of Figures 1-8 (SEQ ID NOS:25-32) may be (i) one in which one or more of the amino acid residues are substituted with a conserved or non-conserved amino acid residue (preferably a conserved amino acid residue) and such substituted amino acid residue may or may not be one encoded by the genetic code, or (ii) one in which one or more of the amino acid residues includes a substituent group, or (iii) one in which the mature enzyme is fused with another compound, such as a compound to increase the half-life of the enzyme (for example, polyethylene glycol), or (iv) one in which the additional amino acids are fused to the mature enzyme, such as a leader or secretory sequence or a sequence which is employed for purification of the mature enzyme or a proprotein sequence. Such fragments, derivatives and analogs are deemed to be within the scope of those skilled in the art from the teachings herein.

The enzymes and polynucleotides of the present invention are preferably provided in an isolated form, and preferably are purified to homogeneity.

The term "isolated" means that the material is removed from its original environment (e.g., the natural environment if it is naturally occurring). For example, a naturally-occurring polynucleotide or enzyme present in a living animal is not isolated, but the same polynucleotide or enzyme, separated from some or all of the coexisting materials in the natural system, is isolated. Such polynucleotides could be part of a vector and/or such polynucleotides or enzymes could be part of a composition, and still be isolated in that such vector or composition is not part of its natural environment.

The enzymes of the present invention include the enzymes of SEQ ID NOS:25-32 (in particular the mature enzyme) as well as enzymes which have at least 70% similarity (preferably at least 70% identity) to the enzymes of SEQ ID NOS:25-32 and more preferably at least 90% similarity (more preferably at least 90% identity) to the enzymes of SEQ ID NOS:25-32 and still more preferably at least 95% similarity (still more preferably at least 95% identity) to the enzymes of SEQ ID NOS:25-32 and also include

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portions of such enzymes with such portion of the enzyme generally containing at least 30 amino acids and more preferably at least 50 amino acids.

As known in the art "similarity" between two enzymes is determined by comparing the amino acid sequence and its conserved amino acid substitutes of one enzyme to the sequence of a second enzyme.

A variant, *i.e.* a "fragment", "analog" or "derivative" polypeptide, and reference polypeptide may differ in amino acid sequence by one or more substitutions, additions, deletions, fusions and truncations, which may be present in any combination.

Among preferred variants are those that vary from a reference by conservative amino acid substitutions. Such substitutions are those that substitute a given amino acid in a polypeptide by another amino acid of like characteristics. Typically seen as conservative substitutions are the replacements, one for another, among the aliphatic amino acids Ala, Val, Leu and Ile; interchange of the hydroxyl residues Ser and Thr, exchange of the acidic residues Asp and Glu, substitution between the amide residues Asn and Gln, exchange of the basic residues Lys and Arg and replacements among the aromatic residues Phe, Tyr.

Most highly preferred are variants which retain the same biological function and activity as the reference polypeptide from which it varies.

Fragments or portions of the enzymes of the present invention may be employed for producing the corresponding full-length enzyme by peptide synthesis; therefore, the fragments may be employed as intermediates for producing the full-length enzymes. Fragments or portions of the polynucleotides of the present invention may be used to synthesize full-length polynucleotides of the present invention.

The present invention also relates to vectors which include polynucleotides of the present invention, host cells which are genetically engineered with vectors of the invention and the production of enzymes of the invention by recombinant techniques.

Host cells are genetically engineered (transduced or transformed or transfected) with the vectors of this invention which may be, for example, a cloning vector such as an expression vector. The vector may be, for example, in the form of a plasmid, a phage, *etc.* The engineered host cells can be cultured in conventional nutrient media modified as appropriate for activating promoters, selecting transformants or amplifying the genes of the present invention. The culture conditions, such as temperature, pH and the like, are those previously used with the host cell selected for expression, and will be apparent to the ordinarily skilled artisan.

The polynucleotides of the present invention may be employed for producing enzymes by recombinant techniques. Thus, for example, the polynucleotide may be included in any one of a variety of expression vectors for expressing an enzyme. Such vectors include chromosomal, nonchromosomal and synthetic DNA sequences, *e.g.*, derivatives of SV40; bacterial plasmids; phage DNA; baculovirus; yeast plasmids; vectors derived from combinations of plasmids and phage DNA, viral DNA such as vaccinia, adenovirus, fowl pox virus, and pseudorabies. However, any other vector may be used as long as it is replicable and viable in the host.

The appropriate DNA sequence may be inserted into the vector by a variety of procedures. In general, the DNA sequence is inserted into an appropriate restriction endonuclease site(s) by procedures known in the art. Such procedures and others are deemed to be within the scope of those skilled in the art.

The DNA sequence in the expression vector is operatively linked to an appropriate expression control sequence(s) (promoter) to direct mRNA synthesis. As representative examples of such promoters, there may be mentioned: LTR or SV40

promoter, the *E. coli*. *lac* or *trp*, the phage lambda P_L promoter and other promoters known to control expression of genes in prokaryotic or eukaryotic cells or their viruses. The expression vector also contains a ribosome binding site for translation initiation and a transcription terminator. The vector may also include appropriate sequences for amplifying expression.

In addition, the expression vectors preferably contain one or more selectable marker genes to provide a phenotypic trait for selection of transformed host cells such as dihydrofolate reductase or neomycin resistance for eukaryotic cell culture, or such as tetracycline or ampicillin resistance in *E. coli*.

The vector containing the appropriate DNA sequence as hereinabove described, as well as an appropriate promoter or control sequence, may be employed to transform an appropriate host to permit the host to express the protein.

As representative examples of appropriate hosts, there may be mentioned: bacterial cells, such as *E. coli*, *Streptomyces*, *Bacillus subtilis*; fungal cells, such as yeast; insect cells such as *Drosophila S2* and *Spodoptera Sf9*; animal cells such as CHO, COS or Bowes melanoma; adenoviruses; plant cells, *etc.* The selection of an appropriate host is deemed to be within the scope of those skilled in the art from the teachings herein.

More particularly, the present invention also includes recombinant constructs comprising one or more of the sequences as broadly described above. The constructs comprise a vector, such as a plasmid or viral vector, into which a sequence of the invention has been inserted, in a forward or reverse orientation. In a preferred aspect of this embodiment, the construct further comprises regulatory sequences, including, for example, a promoter, operably linked to the sequence. Large numbers of suitable vectors and promoters are known to those of skill in the art, and are commercially available. The following vectors are provided by way of example; Bacterial: pQE70,

pQE60, pQE-9 (Qiagen), pBluescript II KS, ptrc99a, pKK223-3, pDR540, pRIT2T (Pharmacia); Eukaryotic: pXT1, pSG5 (Stratagene) pSVK3, pBPV, pMSG, pSVL SV40 (Pharmacia). However, any other plasmid or vector may be used as long as they are replicable and viable in the host.

Promoter regions can be selected from any desired gene using CAT (chloramphenicol transferase) vectors or other vectors with selectable markers. Two appropriate vectors are pKK232-8 and pCM7. Particular named bacterial promoters include lacI, lacZ, T3, T7, gpt, lambda P_R, P_L and trp. Eukaryotic promoters include CMV immediate early, HSV thymidine kinase, early and late SV40, LTRs from retrovirus, and mouse metallothionein-I. Selection of the appropriate vector and promoter is well within the level of ordinary skill in the art.

In a further embodiment, the present invention relates to host cells containing the above-described constructs. The host cell can be a higher eukaryotic cell, such as a mammalian cell, or a lower eukaryotic cell, such as a yeast cell, or the host cell can be a prokaryotic cell, such as a bacterial cell. Introduction of the construct into the host cell can be effected by calcium phosphate transfection, DEAE-Dextran mediated transfection, or electroporation (Davis, L., Dibner, M., Battey, I., *Basic Methods in Molecular Biology*, (1986)).

The constructs in host cells can be used in a conventional manner to produce the gene product encoded by the recombinant sequence. Alternatively, the enzymes of the invention can be synthetically produced by conventional peptide synthesizers.

Mature proteins can be expressed in mammalian cells, yeast, bacteria, or other cells under the control of appropriate promoters. Cell-free translation systems can also be employed to produce such proteins using RNAs derived from the DNA constructs of the present invention. Appropriate cloning and expression vectors for use with prokaryotic and eukaryotic hosts are described by Sambrook *et al.*, *Molecular Cloning*:

A Laboratory Manual, Second Edition, Cold Spring Harbor, N.Y., (1989), the disclosure of which is hereby incorporated by reference.

Transcription of the DNA encoding the enzymes of the present invention by higher eukaryotes is increased by inserting an enhancer sequence into the vector. Enhancers are cis-acting elements of DNA, usually about from 10 to 300 bp that act on a promoter to increase its transcription. Examples include the SV40 enhancer on the late side of the replication origin bp 100 to 270, a cytomegalovirus early promoter enhancer, the polyoma enhancer on the late side of the replication origin, and adenovirus enhancers.

Generally, recombinant expression vectors will include origins of replication and selectable markers permitting transformation of the host cell, *e.g.*, the ampicillin resistance gene of *E. coli* and *S. cerevisiae* TRP1 gene, and a promoter derived from a highly-expressed gene to direct transcription of a downstream structural sequence. Such promoters can be derived from operons encoding glycolytic enzymes such as 3-phosphoglycerate kinase (PGK), α -factor, acid phosphatase, or heat shock proteins, among others. The heterologous structural sequence is assembled in appropriate phase with translation initiation and termination sequences, and preferably, a leader sequence capable of directing secretion of translated enzyme. Optionally, the heterologous sequence can encode a fusion enzyme including an N-terminal identification peptide imparting desired characteristics, *e.g.*, stabilization or simplified purification of expressed recombinant product.

Useful expression vectors for bacterial use are constructed by inserting a structural DNA sequence encoding a desired protein together with suitable translation initiation and termination signals in operable reading phase with a functional promoter. The vector will comprise one or more phenotypic selectable markers and an origin of replication to ensure maintenance of the vector and to, if desirable, provide amplification within the host. Suitable prokaryotic hosts for transformation include *E.*

coli, *Bacillus subtilis*, *Salmonella typhimurium* and various species within the genera *Pseudomonas*, *Streptomyces*, and *Staphylococcus*, although others may also be employed as a matter of choice.

As a representative but nonlimiting example, useful expression vectors for bacterial use can comprise a selectable marker and bacterial origin of replication derived from commercially available plasmids comprising genetic elements of the well known cloning vector pBR322 (ATCC 37017). Such commercial vectors include, for example, pKK223-3 (Pharmacia Fine Chemicals, Uppsala, Sweden) and pGEM1 (Promega Biotec, Madison, WI, USA). These pBR322 "backbone" sections are combined with an appropriate promoter and the structural sequence to be expressed.

Following transformation of a suitable host strain and growth of the host strain to an appropriate cell density, the selected promoter is induced by appropriate means (*e.g.*, temperature shift or chemical induction) and cells are cultured for an additional period.

Cells are typically harvested by centrifugation, disrupted by physical or chemical means, and the resulting crude extract retained for further purification.

Microbial cells employed in expression of proteins can be disrupted by any convenient method, including freeze-thaw cycling, sonication, mechanical disruption, or use of cell lysing agents, such methods are well known to those skilled in the art.

Various mammalian cell culture systems can also be employed to express recombinant protein. Examples of mammalian expression systems include the COS-7 lines of monkey kidney fibroblasts, described by Gluzman, *Cell*, 23:175 (1981), and other cell lines capable of expressing a compatible vector, for example, the C127, 3T3, CHO, HeLa and BHK cell lines. Mammalian expression vectors will comprise an origin of replication, a suitable promoter and enhancer, and also any necessary ribosome

binding sites, polyadenylation site, splice donor and acceptor sites, transcriptional termination sequences, and 5' flanking nontranscribed sequences. DNA sequences derived from the SV40 splice, and polyadenylation sites may be used to provide the required nontranscribed genetic elements.

The enzyme can be recovered and purified from recombinant cell cultures by methods including ammonium sulfate or ethanol precipitation, acid extraction, anion or cation exchange chromatography, phosphocellulose chromatography, hydrophobic interaction chromatography, affinity chromatography, hydroxylapatite chromatography and lectin chromatography. Protein refolding steps can be used, as necessary, in completing configuration of the mature protein. Finally, high performance liquid chromatography (HPLC) can be employed for final purification steps.

The enzymes of the present invention may be a naturally purified product, or a product of chemical synthetic procedures, or produced by recombinant techniques from a prokaryotic or eukaryotic host (for example, by bacterial, yeast, higher plant, insect and mammalian cells in culture). Depending upon the host employed in a recombinant production procedure, the enzymes of the present invention may be glycosylated or may be non-glycosylated. Enzymes of the invention may or may not also include an initial methionine amino acid residue.

Transaminases are a group of key enzymes in the metabolism of amino acids and amino sugars and are found in all organisms from microbes to mammals. In the transamination reaction, an amino group is transferred from an amino acid to an α -keto acid. Pyridoxal phosphate is required as a co-factor to mediate the transfer of the amino group without liberation of ammonia.

Amino acids currently have applications as additives to animal feed, human nutritional supplements, components in infusion solutions, and synthetic intermediates for manufacture of pharmaceuticals and agricultural products. For example, L-glutamic

acid is best known as a flavor enhancer for human food. L-lysine and L-methionine are large volume additives to animal feed and human supplements. L-tryptophan and L-threonine have similar potential applications. L-phenylalanine and L-aspartic acid have very important market potential as key components in the manufacture of the low-calorie sweetener aspartame, and other promising low-calorie sweeteners have compositions containing certain amino acids as well. Infusion solutions require a large range of amino acids including those essential ones in human diets.

Transaminases are highly stereoselective, and most use L-amino acids as substrates. Using the approach disclosed in a commonly assigned, copending provisional application Serial No. 60/008,316, filed on December 7, 1995 and entitled "Combinatorial Enzyme Development," the disclosure of which is incorporated herein by reference in its entirety, one can convert the transaminases of the invention to use D-amino acids as substrates. Such conversion makes possible a broader array of transaminase applications. For instance, D-valine can be used in the manufacture of synthetic pyrethroids. D-phenylglycine and its derivatives can be useful as components of β -lactam antibiotics. Further, the thermostable transaminases have superior stability at higher temperatures and in organic solvents. Thus, they are better suited to utilize either L- and/or D-amino acids for production of optically pure chiral compounds used in pharmaceutical, agricultural, and other chemical manufactures.

There are a number of reasons to employ transaminases in industrial-scale production of amino acids and their derivatives.

- 1) Transaminases can catalyze stereoselective synthesis of D- or L-amino acids from their corresponding α -keto acids. Therefore no L- or D-isomers are produced, and no resolution is required.

- 2) Transaminases have uniformly high catalytic rates, capable of converting up to 400 μ moles of substrates per minute per mg enzyme.

3) Many required α -keto acids can be conveniently prepared by chemical synthesis at low cost.

4) The capital investment for an immobilized enzyme process using transaminases is much lower than for a large scale fermentation process, and productivity of the bioreactor is often an order of magnitude higher.

5) The technology is generally applicable to a broad range of D- or L-amino acids because transaminases exist with varying specificities. Such broad scope allows a number of different L- or D-amino acids to be produced with the same equipment and often the same biocatalyst.

Antibodies generated against the enzymes corresponding to a sequence of the present invention can be obtained by direct injection of the enzymes into an animal or by administering the enzymes to an animal, preferably a nonhuman. The antibody so obtained will then bind the enzymes itself. In this manner, even a sequence encoding only a fragment of the enzymes can be used to generate antibodies binding the whole native enzymes. Such antibodies can then be used to isolate the enzyme from cells expressing that enzyme.

For preparation of monoclonal antibodies, any technique which provides antibodies produced by continuous cell line cultures can be used. Examples include the hybridoma technique (Kohler and Milstein, *Nature*, 256:495-497, 1975), the trioma technique, the human B-cell hybridoma technique (Kozbor *et al.*, *Immunology Today* 4:72, 1983), and the EBV-hybridoma technique to produce human monoclonal antibodies (Cole *et al.*, in *Monoclonal Antibodies and Cancer Therapy*, Alan R. Liss, Inc., pp. 77-96, 1985).

Techniques described for the production of single chain antibodies (U.S. Patent 4,946,778) can be adapted to produce single chain antibodies to immunogenic enzyme

products of this invention. Also, transgenic mice may be used to express humanized antibodies to immunogenic enzyme products of this invention.

Antibodies generated against an enzyme of the present invention may be used in screening for similar enzymes from other organisms and samples. Such screening techniques are known in the art, for example, one such screening assay is described in Sambrook and Maniatis, *Molecular Cloning: A Laboratory Manual* (2d Ed.), vol. 2:Section 8.49, Cold Spring Harbor Laboratory, 1989, which is hereby incorporated by reference in its entirety.

The present invention will be further described with reference to the following examples; however, it is to be understood that the present invention is not limited to such examples. All parts or amounts, unless otherwise specified, are by weight.

In order to facilitate understanding of the following examples certain frequently occurring methods and/or terms will be described.

"Plasmids" are designated by a lower case "p" preceded and/or followed by capital letters and/or numbers. The starting plasmids herein are either commercially available, publicly available on an unrestricted basis, or can be constructed from available plasmids in accord with published procedures. In addition, equivalent plasmids to those described are known in the art and will be apparent to the ordinarily skilled artisan.

"Digestion" of DNA refers to catalytic cleavage of the DNA with a restriction enzyme that acts only at certain sequences in the DNA. The various restriction enzymes used herein are commercially available and their reaction conditions, cofactors and other requirements were used as would be known to the ordinarily skilled artisan. For analytical purposes, typically 1 μ g of plasmid or DNA fragment is used with about 2 units of enzyme in about 20 μ l of buffer solution. For the purpose of isolating DNA

fragments for plasmid construction, typically 5 to 50 μg of DNA are digested with 20 to 250 units of enzyme in a larger volume. Appropriate buffers and substrate amounts for particular restriction enzymes are specified by the manufacturer. Incubation times of about 1 hour at 37°C are ordinarily used, but may vary in accordance with the supplier's instructions. After digestion the reaction is electrophoresed directly on a polyacrylamide gel to isolate the desired fragment.

Size separation of the cleaved fragments is performed using 8 percent polyacrylamide gel described by Goeddel *et al.*, *Nucleic Acids Res.*, 8:4057 (1980).

"Oligonucleotides" refers to either a single stranded polydeoxynucleotide or two complementary polydeoxynucleotide strands which may be chemically synthesized. Such synthetic oligonucleotides have no 5' phosphate and thus will not ligate to another oligonucleotide without adding a phosphate with an ATP in the presence of a kinase. A synthetic oligonucleotide will ligate to a fragment that has not been dephosphorylated.

"Ligation" refers to the process of forming phosphodiester bonds between two double stranded nucleic acid fragments (Maniatis, T., *et al.*, *Id.*, p. 146). Unless otherwise provided, ligation may be accomplished using known buffers and conditions with 10 units of T4 DNA ligase ("ligase") per 0.5 μg of approximately equimolar amounts of the DNA fragments to be ligated.

Unless otherwise stated, transformation was performed as described in Sambrook and Maniatis, *Molecular Cloning: A Laboratory Manual*, Cold Spring Harbor Laboratory, 1989.

Example 1

Bacterial Expression and Purification of Transaminases and Aminotransferases

DNA encoding the enzymes of the present invention, SEQ ID NOS:25 through 32, were initially amplified from a pBluescript vector containing the DNA by the PCR technique using the primers noted herein. The amplified sequences were then inserted into the respective PQE vector listed beneath the primer sequences, and the enzyme was expressed according to the protocols set forth herein. The genomic DNA has also been used as a template for the PCR amplification, *i.e.*, once a positive clone has been identified and primer sequences determined using the cDNA, it was then possible to return to the genomic DNA and directly amplify the desired sequence(s) there. The 5' and 3' primer sequences and the vector for the respective genes are as follows:

25-32
37,32

Aquifex Aspartate Transaminase A

aspa501 5' CCGAGAATTCATTAAAGAGGAGAAATTA ACTATGATTGAAGACCCTATGGAC (SEQ. ID NO:1)

aspa301 3' CGAAGATCTTTAGCACTTCTCTCAGGTTC (SEQ. ID NO:2)

vector: pQET1

Aquifex Aspartate Aminotransferase B

aspb501 5' CCGAGAATTCATTAAAGAGGAGAAATTA ACTATGGACAGGCTTGAAAAAGTA (SEQ ID NO:3)

aspb301 3' CGGAAGATCTTCAGCTAAGCTTCTCTAAGAA (SEQ ID NO:4)

vector: pQET1

Aquifex Adenosyl-8-amino-7-oxononanoate Aminotransferase

ameth501 5' CCGACAATTGATTAAAGAGGAGAAATTA ACTATGTGGGAATTAGACCCTAAA (SEQ ID NO:5)

ameth301 3' CGGAGGATCCCTACACCTCTTTTCAAGCT (SEQ ID NO:6)

vector: pQET12

Aquifex Acetylornithine Aminotransferase

aorn 501 5' CCGACAATTGATTAAAGAGGAGAAATTA ACTATGACATACTTAATGAACAAT (SEQ ID NO:7)

aorn 301 3' CGGAAGATCTTTATGAGAAGTCCCTTTCAAG (SEQ ID NO:8)

vector: pQET12

Ammonifex degensii Aspartate Aminotransferase

adasp 501 5' CCGAGAATTCATTAAAGAGGAGAAATTAAGTATGCGGAACTGGCCGAGCGG (SEQ ID NO:9)

adasp 301 3' CGGAGGATCCTTAAAGTGCCGCTTCGATCAA (SEQ ID NO:10)

vector: pQET12

Aquifex Glucosamine:Fructose-6-phosphate Aminotransferase

glut 501 5' CCGACAATTGATTAAAGAGGAGAAATTAAGTATGTGCGGGATAGTCGGATAC (SEQ ID NO:11)

glut 301 3' CGGAAGATCTTTATTCCACCGTGACCGTTTT (SEQ ID NO:12)

vector: pQET1

Aquifex Histadine-phosphate Aminotransferase

his 501 5' CCGACAATTGATTAAAGAGGAGAAATTAAGTATGATACCCAGAGGATTAAG (SEQ ID NO:13)

his 301 3' CGGAAGATCTTTAAAGAGAGCTTGAAAGGGA (SEQ ID NO:14)

vector: pQET1

Pyrobaculum aerophilum Branched Chain Aminotransferase

bcat 501 5' CCGAGAATTCATTAAAGAGGAGAAATTAAGTATGAAGCCGTACGCTAAATAT (SEQ ID NO:15)

bcat 301 3' CGGAAGATCTCTAATACACAGGAGTGATCCA (SEQ ID NO:16)

vector: pQET1

Ammonifex degensii hp aminotransferase

5' -CCGAGAATTCATTAAAGAGGAGAAATTAAGTATGGCAGTCAAAGTGCGGCCT 31

3' -CGGAGGATCCTTATCCAAAGCTTCCAGGAAG 34

Homology information:

Closest to *Bacillus subtilis* (reference: Henner D.J., Band L., Flaggs G., Chen E.; 20

Gene 49:147-152(1986). Percent Similarity: 65.084 Percent Identity: 44.134 20

Aquifex aspartate aminotransferase

5' CCGAGAATTCATTAAAGAGGAGAAATTAAGTATGAGAAAAGGACTTGCAAGT 37

3' CGGAGGATCCTTAGATCTCTTCAAGGGCTTT 38

Closest to *Bacillus subtilis* (Sorokin, A.V., Azevedo, V., Zumstein, E., Galleron, N., Ehrlich, S.D. and Serror, P. Determination and analysis of the nucleotide sequence of the *Bacillus subtilis* chromosome region between *serA* and *kdg* loci cloned in yeast artificial chromosome Unpublished (1995). Percent Similarity: 71.611 Percent Identity: 52.685

The restriction enzyme sites indicated correspond to the restriction enzyme sites on the bacterial expression vector indicated for the respective gene (Qiagen, Inc. Chatsworth, CA). The pQE vector encodes antibiotic resistance (Amp^r), a bacterial origin of replication (ori), an IPTG-regulatable promoter operator (P/O), a ribosome binding site (RBS), a 6-His tag and restriction enzyme sites.

The pQE vector was digested with the restriction enzymes indicated. The amplified sequences were ligated into the respective pQE vector and inserted in frame with the sequence encoding for the RBS. The ligation mixture was then used to transform the *E. coli* strain M15/pREP4 (Qiagen, Inc.) by electroporation. M15/pREP4 contains multiple copies of the plasmid pREP4, which expresses the lacI repressor and also confers kanamycin resistance (Kan^r). Transformants were identified by their ability to grow on LB plates and ampicillin/kanamycin resistant colonies were selected. Plasmid DNA was isolated and confirmed by restriction analysis. Clones containing the desired constructs were grown overnight (O/N) in liquid culture in LB media supplemented with both Amp (100 ug/ml) and Kan (25 ug/ml). The O/N culture was used to inoculate a large culture at a ratio of 1:100 to 1:250. The cells were grown to an optical density 600 (O.D.⁶⁰⁰) of between 0.4 and 0.6. IPTG ("Isopropyl-B-D-thiogalacto pyranoside") was then added to a final concentration of 1 mM. IPTG induces by inactivating the lacI repressor, clearing the P/O leading to increased gene expression. Cells were grown an extra 3 to 4 hours. Cells were then harvested by centrifugation.

The primer sequences set out above may also be employed to isolate the target gene from the deposited material by hybridization techniques described above.

Example 2

Isolation of a Selected Clone from the Deposited Genomic Clones

The two oligonucleotide primers corresponding to the gene of interest are used to amplify the gene from the deposited material. A polymerase chain reaction is carried out in 25 μ l of reaction mixture with 0.1 μ g of the DNA of the gene of interest. The reaction mixture is 1.5-5 mM MgCl₂, 0.01% (w/v) gelatin, 20 μ M each of dATP, dCTP, dGTP, dTTP, 25 pmol of each primer and 1.25 Unit of Taq polymerase. Thirty cycles of PCR (denaturation at 94°C for 1 min; annealing at 55°C for 1 min; elongation at 72°C for 1 min) are performed with the Perkin-Elmer Cetus 9600 thermal cycler. The amplified product is analyzed by agarose gel electrophoresis and the DNA band with expected molecular weight is excised and purified. The PCR product is verified to be the gene of interest by subcloning and sequencing the DNA product.

Numerous modifications and variations of the present invention are possible in light of the above teachings and, therefore, within the scope of the appended claims, the invention may be practiced otherwise than as particularly described.

03385537 090299

SEQUENCE LISTING

(1) GENERAL INFORMATION:

(i) APPLICANTS:

WARREN, Patrick V.

SWANSON, Ronald V.

(ii) TITLE OF INVENTION:

TRANSAMINASES AND AMINOTRANSFERASES

(iii) NUMBER OF SEQUENCES: 40

(iv) CORRESPONDENCE ADDRESS:

(A) ADDRESSEE: CARELLA, BYRNE, BAIN, GILFILLAN,
CECCHI, STEWART & OLSTEIN

(B) STREET: 6 BECKER FARM ROAD

(C) CITY: ROSELAND

(D) STATE: NEW JERSEY

(E) COUNTRY: USA

(F) ZIP: 07068

(v) COMPUTER READABLE FORM:

(A) MEDIUM TYPE: 3.5 INCH DISKETTE

(B) COMPUTER: IBM PS/2

(C) OPERATING SYSTEM: MS-DOS

(D) SOFTWARE: WORD PERFECT 5.1

(vi) CURRENT APPLICATION DATA:

(A) APPLICATION NUMBER: Unassigned

(B) FILING DATE: Concurrently

(C) CLASSIFICATION: Unassigned

(vii) PRIOR APPLICATION DATA:

(A) APPLICATION NUMBER:

(B) FILING DATE:

(C) CLASSIFICATION:

(viii) ATTORNEY/AGENT INFORMATION:

(A) NAME: HERRON, CHARLES J.

(B) REGISTRATION NUMBER: 28,019

(C) REFERENCE/DOCKET NUMBER: 331400-38

(ix) TELECOMMUNICATION INFORMATION:

(A) TELEPHONE: 201-994-1700

(B) TELEFAX: 201-994-1744

(2) INFORMATION FOR SEQ ID NO:1:

(i) SEQUENCE CHARACTERISTICS

(A) LENGTH: 52 NUCLEOTIDES

(B) TYPE: NUCLEIC ACID

(C) STRANDEDNESS: SINGLE

(D) TOPOLOGY: LINEAR

(ii) MOLECULE TYPE: cDNA

(xi) SEQUENCE DESCRIPTION: SEQ ID NO:1:

CCGAGAATTC ATTAAAGAGG AGAAATTAAC TATGATTGAA GACCCTATGG AC

52

(2) INFORMATION FOR SEQ ID NO:2:

(i) SEQUENCE CHARACTERISTICS

- (A) LENGTH: 31 NUCLEOTIDES
- (B) TYPE: NUCLEIC ACID
- (C) STRANDEDNESS: SINGLE
- (D) TOPOLOGY: LINEAR

(ii) MOLECULE TYPE: cDNA

(xi) SEQUENCE DESCRIPTION: SEQ ID NO:2:

CGGAAGATCT TTAAGCACTT CTCTCAGGTT C

31

(2) INFORMATION FOR SEQ ID NO:3:

(i) SEQUENCE CHARACTERISTICS

- (A) LENGTH: 52 NUCLEOTIDES
- (B) TYPE: NUCLEIC ACID
- (C) STRANDEDNESS: SINGLE
- (D) TOPOLOGY: LINEAR

(ii) MOLECULE TYPE: cDNA

(xi) SEQUENCE DESCRIPTION: SEQ ID NO:3:

CCGAGAATTC ATTAAAGAGG AGAAATTAAC TATGGACAGG CTTGAAAAAG TA

52

(2) INFORMATION FOR SEQ ID NO:4:

(i) SEQUENCE CHARACTERISTICS

- (A) LENGTH: 31 NUCLEOTIDES
- (B) TYPE: NUCLEIC ACID
- (C) STRANDEDNESS: SINGLE
- (D) TOPOLOGY: LINEAR

(ii) MOLECULE TYPE: cDNA

(xi) SEQUENCE DESCRIPTION: SEQ ID NO:4:

CGGAAGATCT TCAGCTAAGC TTCTCTAAGA A

31

(2) INFORMATION FOR SEQ ID NO:5:

(i) SEQUENCE CHARACTERISTICS

- (A) LENGTH: 52 NUCLEOTIDES
- (B) TYPE: NUCLEIC ACID
- (C) STRANDEDNESS: SINGLE
- (D) TOPOLOGY: LINEAR

(ii) MOLECULE TYPE: cDNA

(xi) SEQUENCE DESCRIPTION: SEQ ID NO:5:

CCGACAATTG ATTAAAGAGG AGAAATTAAC TATGTGGGAA TTAGACCCTA AA

52

(2) INFORMATION FOR SEQ ID NO:6:

- (i) SEQUENCE CHARACTERISTICS
 - (A) LENGTH: 31 NUCLEOTIDES
 - (B) TYPE: NUCLEIC ACID
 - (C) STRANDEDNESS: SINGLE
 - (D) TOPOLOGY: LINEAR

(ii) MOLECULE TYPE: cDNA

(xi) SEQUENCE DESCRIPTION: SEQ ID NO:6:

CGGAGGATCC CTACACCTGT TTTTCAAGCT C

31

(2) INFORMATION FOR SEQ ID NO:7:

- (i) SEQUENCE CHARACTERISTICS
 - (A) LENGTH: 52 NUCLEOTIDES
 - (B) TYPE: NUCLEIC ACID
 - (C) STRANDEDNESS: SINGLE
 - (D) TOPOLOGY: LINEAR

(ii) MOLECULE TYPE: cDNA

(xi) SEQUENCE DESCRIPTION: SEQ ID NO:7:

CCGACAATTG ATTAAAGAGG AGAAATTAAC TATGACATAC TTAATGAACA AT

52

(2) INFORMATION FOR SEQ ID NO:8:

- (i) SEQUENCE CHARACTERISTICS
 - (A) LENGTH: 31 NUCLEOTIDES
 - (B) TYPE: NUCLEIC ACID
 - (C) STRANDEDNESS: SINGLE
 - (D) TOPOLOGY: LINEAR

(ii) MOLECULE TYPE: cDNA

(xi) SEQUENCE DESCRIPTION: SEQ ID NO:8:

CGGAAGATCT TTATGAGAAG TCCCTTTCAA G

31

(2) INFORMATION FOR SEQ ID NO:9:

- (i) SEQUENCE CHARACTERISTICS
 - (A) LENGTH: 52 NUCLEOTIDES
 - (B) TYPE: NUCLEIC ACID
 - (C) STRANDEDNESS: SINGLE
 - (D) TOPOLOGY: LINEAR

(ii) MOLECULE TYPE: cDNA

(xi) SEQUENCE DESCRIPTION: SEQ ID NO:9:

CCGAGAATTC ATTAAAGAGG AGAAATTAAC TATGCGGAAA CTGGCCGAGC GG

52

(2) INFORMATION FOR SEQ ID NO:10:

- (i) SEQUENCE CHARACTERISTICS
 - (A) LENGTH: 31 NUCLEOTIDES
 - (B) TYPE: NUCLEIC ACID
 - (C) STRANDEDNESS: SINGLE
 - (D) TOPOLOGY: LINEAR
- (ii) MOLECULE TYPE: cDNA
- (xi) SEQUENCE DESCRIPTION: SEQ ID NO:10:

CGGAGGATCC TTAAAGTGCC GCTTCGATCA A

31

- (2) INFORMATION FOR SEQ ID NO:11:
 - (i) SEQUENCE CHARACTERISTICS
 - (A) LENGTH: 52 NUCLEOTIDES
 - (B) TYPE: NUCLEIC ACID
 - (C) STRANDEDNESS: SINGLE
 - (D) TOPOLOGY: LINEAR
 - (ii) MOLECULE TYPE: cDNA
 - (xi) SEQUENCE DESCRIPTION: SEQ ID NO:11:

CCGACAATTG ATTAAAGAGG AGAAATTAAC TATGTGCGGG ATAGTCGGAT AC

52

- (2) INFORMATION FOR SEQ ID NO:12:
 - (i) SEQUENCE CHARACTERISTICS
 - (A) LENGTH: 31 NUCLEOTIDES
 - (B) TYPE: NUCLEIC ACID
 - (C) STRANDEDNESS: SINGLE
 - (D) TOPOLOGY: LINEAR
 - (ii) MOLECULE TYPE: cDNA
 - (xi) SEQUENCE DESCRIPTION: SEQ ID NO:12:

CGGAAGATCT TTATTCCACC GTGACCGTTT T

31

- (2) INFORMATION FOR SEQ ID NO:13:
 - (i) SEQUENCE CHARACTERISTICS
 - (A) LENGTH: 52 NUCLEOTIDES
 - (B) TYPE: NUCLEIC ACID
 - (C) STRANDEDNESS: SINGLE
 - (D) TOPOLOGY: LINEAR
 - (ii) MOLECULE TYPE: cDNA
 - (xi) SEQUENCE DESCRIPTION: SEQ ID NO:13:

CCGACAATTG ATTAAAGAGG AGAAATTAAC TATGATACCC CAGAGGATTA AG

52

- (2) INFORMATION FOR SEQ ID NO:14:
 - (i) SEQUENCE CHARACTERISTICS
 - (A) LENGTH: 31 NUCLEOTIDES
 - (B) TYPE: NUCLEIC ACID
 - (C) STRANDEDNESS: SINGLE
 - (D) TOPOLOGY: LINEAR

(ii) MOLECULE TYPE: cDNA

(xi) SEQUENCE DESCRIPTION: SEQ ID NO:14:

CGGAAGATCT TTAAAGAGAG CTTGAAAGGG A

31

(2) INFORMATION FOR SEQ ID NO:15:

(i) SEQUENCE CHARACTERISTICS

(A) LENGTH: 52 NUCLEOTIDES

(B) TYPE: NUCLEIC ACID

(C) STRANDEDNESS: SINGLE

(D) TOPOLOGY: LINEAR

(ii) MOLECULE TYPE: cDNA

(xi) SEQUENCE DESCRIPTION: SEQ ID NO:15:

CCGAGAATTC ATTAAAGAGG AGAAATTAAC TATGAAGCCG TACGCTAAAT AT

52

(2) INFORMATION FOR SEQ ID NO:16:

(i) SEQUENCE CHARACTERISTICS

(A) LENGTH: 31 NUCLEOTIDES

(B) TYPE: NUCLEIC ACID

(C) STRANDEDNESS: SINGLE

(D) TOPOLOGY: LINEAR

(ii) MOLECULE TYPE: cDNA

(xi) SEQUENCE DESCRIPTION: SEQ ID NO:16:

CGGAAGATCT CTAATACACA GGAGTGATCC A

31

(2) INFORMATION FOR SEQ ID NO:17:

(i) SEQUENCE CHARACTERISTICS

(A) LENGTH: 1245 NUCLEOTIDES

(B) TYPE: NUCLEIC ACID

(C) STRANDEDNESS: SINGLE

(D) TOPOLOGY: LINEAR

(ii) MOLECULE TYPE: GENOMIC DNA

(xi) SEQUENCE DESCRIPTION: SEQ ID NO:17:

ATG ATT GAA GAC CCT ATG GAC TGG GCT TTT CCG AGG ATA AAG AGA CTG
Met Ile Glu Asp Pro Met Asp Trp Ala Phe Pro Arg Ile Lys Arg Leu
5 10 15

48

CCT CAG TAT GTC TTC TCT CTC GTT AAC GAA CTC AAG TAC AAG CTA AGG
Pro Gln Tyr Val Phe Ser Leu Val Asn Glu Leu Lys Tyr Lys Leu Arg
20 25 30

96

CGT GAA GGC GAA GAT GTA GTG GAT CTT GGT ATG GGC AAT CCT AAC ATG
Arg Glu Gly Glu Asp Val Val Asp Leu Gly Met Gly Asn Pro Asn Met
35 40 45

144

CCT CCA GCA AAG CAC ATA ATA GAT AAA CTC TGC GAA GTG GCT CAA AAG
Pro Pro Ala Lys His Ile Ile Asp Lys Leu Cys Glu Val Ala Gln Lys
50 55 60

192

CCG AAC GTT CAC GGA TAT TCT GCG TCA AGG GGC ATA CCA AGA CTG AGA Pro Asn Val His Gly Tyr Ser Ala Ser Arg Gly Ile Pro Arg Leu Arg 65 70 75 80	240
AAG GCT ATA TGT AAC TTC TAC GAA GAA AGG TAC GGA GTG AAA CTC GAC Lys Ala Ile Cys Asn Phe Tyr Glu Glu Arg Tyr Gly Val Lys Leu Asp 85 90 95	288
CCT GAG AGG GAG GCT ATA CTA ACA ATC GGT GCA AAG GAA GGG TAT TCT Pro Glu Arg Glu Ala Ile Leu Thr Ile Gly Ala Lys Glu Gly Tyr Ser 100 105 110	336
CAT TTG ATG CTT GCG ATG ATA TCT CCG GGT GAT ACG GTA ATA GTT CCT His Leu Met Leu Ala Met Ile Ser Pro Gly Asp Thr Val Ile Val Pro 115 120 125	384
AAT CCC ACC TAT CCT ATT CAC TAT TAC GCT CCC ATA ATT GCA GGA GGG Asn Pro Thr Tyr Pro Ile His Tyr Tyr Ala Pro Ile Ile Ala Gly Gly 130 135 140	432
GAA GTT CAC TCA ATA CCC CTT AAC TTC TCG GAC GAT CAA GAT CAT CAG Glu Val His Ser Ile Pro Leu Asn Phe Ser Asp Asp Gln Asp His Gln 145 150 155 160	480
GAA GAG TTT TTA AGG AGG CTT TAC GAG ATA GTA AAA ACC GCG ATG CCA Glu Glu Phe Leu Arg Arg Leu Tyr Glu Ile Val Lys Thr Ala Met Pro 165 170 175	528
AAA CCC AAG GCT GTC GTC ATA AGC TTT CCT CAC AAT CCA ACG ACC ATA Lys Pro Lys Ala Val Val Ile Ser Phe Pro His Asn Pro Thr Thr Ile 180 185 190	576
ACG GTA GAA AAG GAC TTT TTT AAA GAA ATA GTT AAG TTT GCA AAG GAA Thr Val Glu Lys Asp Phe Phe Lys Glu Ile Val Lys Phe Ala Lys Glu 195 200 205	624
CAC GGT CTC TGG ATA ATA CAC GAT TTT GCG TAT GCG GAT ATA GCC TTT His Gly Leu Trp Ile Ile His Asp Phe Ala Tyr Ala Asp Ile Ala Phe 210 215 220	672
GAC GGT TAC AAG CCC CCC TCA ATA CTC GAA ATA GAA GGT GCT AAA GAC Asp Gly Tyr Lys Pro Pro Ser Ile Leu Glu Ile Glu Gly Ala Lys Asp 225 230 235 240	720
GTT GCG GTT GAG CTC TAC TCC ATG TCA AAG GGC TTT TCA ATG GCG GGC Val Ala Val Glu Leu Tyr Ser Met Ser Lys Gly Phe Ser Met Ala Gly 245 250 255	768
TGG AGG GTA GCC TTT GTC GTT GGA AAC GAA ATA CTC ATA AAA AAC CTT Trp Arg Val Ala Phe Val Val Gly Asn Glu Ile Leu Ile Lys Asn Leu 260 265 270	816
GCA CAC CTC AAA AGC TAC TTG GAT TAC GGT ATA TTT ACT CCC ATA CAG Ala His Leu Lys Ser Tyr Leu Asp Tyr Gly Ile Phe Thr Pro Ile Gln 275 280 285	864
GTG GCC TCT ATT ATC GCA TTA GAG AGC CCC TAC GAA ATC GTG GAA AAA Val Ala Ser Ile Ile Ala Leu Glu Ser Pro Tyr Glu Ile Val Glu Lys 290 295 300	912
ACC GCA AAG GTT TAC CAA AAA AGA AGA GAC GTT CTG GTG GAA GGG TTA	960

Thr	Ala	Lys	Val	Tyr	Gln	Lys	Arg	Arg	Asp	Val	Leu	Val	Glu	Gly	Leu	
305					310					315					320	
AAC	AGG	CTC	GGC	TGG	AAA	GTA	AAA	AAA	CCT	AAG	GCT	ACC	ATG	TTC	GTC	1008
Asn	Arg	Leu	Gly	Trp	Lys	Val	Lys	Lys	Pro	Lys	Ala	Thr	Met	Phe	Val	
			325						330				335			
TGG	GCA	AAG	ATT	CCC	GAA	TGG	ATA	AAT	ATG	AAC	TCT	CTG	GAC	TTT	TCC	1056
Trp	Ala	Lys	Ile	Pro	Glu	Trp	Ile	Asn	Met	Asn	Ser	Leu	Asp	Phe	Ser	
			340					345					350			
TTG	TTC	CTC	CTA	AAA	GAG	GCG	AAG	GTT	GCG	GTA	TCC	CCG	GGT	GTG	GGC	1104
Leu	Phe	Leu	Leu	Lys	Glu	Ala	Lys	Val	Ala	Val	Ser	Pro	Gly	Val	Gly	
		355					360					365				
TTT	GGT	CAG	TAC	GGA	GAG	GGG	TAC	GTA	AGG	TTT	GCA	CTT	GTA	GAA	AAT	1152
Phe	Gly	Gln	Tyr	Gly	Glu	Gly	Tyr	Val	Arg	Phe	Ala	Leu	Val	Glu	Asn	
	370					375					380					
GAA	CAC	AGG	ATC	AGA	CAG	GCT	ATA	AGG	GGA	ATA	AGG	AAA	GCC	TTC	AGA	1200
Glu	His	Arg	Ile	Arg	Gln	Ala	Ile	Arg	Gly	Ile	Arg	Lys	Ala	Phe	Arg	
385					390					395					400	
AAA	CTC	CAG	AAG	GAG	AGG	AAA	CTT	GAA	CCT	GAG	AGA	AGT	GCT	TAA		1245
Lys	Leu	Gln	Lys	Glu	Arg	Lys	Leu	Glu	Pro	Glu	Arg	Ser	Ala	End		
				405					410				414			

(2) INFORMATION FOR SEQ ID NO:18:

- (i) SEQUENCE CHARACTERISTICS
 (A) LENGTH: 1122 NUCLEOTIDES
 (B) TYPE: NUCLEIC ACID
 (C) STRANDEDNESS: SINGLE
 (D) TOPOLOGY: LINEAR

(ii) MOLECULE TYPE: GENOMIC DNA

(xi) SEQUENCE DESCRIPTION: SEQ ID NO:18:

ATG	GAC	AGG	CTT	GAA	AAA	GTA	TCA	CCC	TTC	ATA	GTA	ATG	GAT	ATC	CTA	48
Met	Asp	Arg	Leu	Glu	Lys	Val	Ser	Pro	Phe	Ile	Val	Met	Asp	Ile	Leu	
			5					10						15		
GCT	CAG	GCC	CAG	AAG	TAC	GAA	GAC	GTA	GTA	CAC	ATG	GAG	ATA	GGA	GAG	96
Ala	Gln	Ala	Gln	Lys	Tyr	Glu	Asp	Val	Val	His	Met	Glu	Ile	Gly	Glu	
			20					25					30			
CCC	GAT	TTA	GAA	CCG	TCT	CCC	AAG	GTA	ATG	GAA	GCT	CTG	GAA	CGT	GCG	144
Pro	Asp	Leu	Glu	Pro	Ser	Pro	Lys	Val	Met	Glu	Ala	Leu	Glu	Arg	Ala	
		35					40					45				
GTG	AAG	GAA	AAG	ACG	TTC	TTC	TAC	ACC	CCT	GCT	CTG	GGA	CTC	TGG	GAA	192
Val	Lys	Glu	Lys	Thr	Phe	Phe	Tyr	Thr	Pro	Ala	Leu	Gly	Leu	Trp	Glu	
	50					55					60					
CTC	AGG	GAA	AGG	ATA	TCG	GAG	TTT	TAC	AGG	AAA	AAG	TAC	AGC	GTT	GAA	240
Leu	Arg	Glu	Arg	Ile	Ser	Glu	Phe	Tyr	Arg	Lys	Lys	Tyr	Ser	Val	Glu	
65					70					75					80	
GTT	TCT	CCA	GAG	AGA	GTC	ATC	GTA	ACT	ACC	GGA	ACT	TCG	GGA	GCG	TTT	288
Val	Ser	Pro	Glu	Arg	Val	Ile	Val	Thr	Thr	Gly	Thr	Ser	Gly	Ala	Phe	
				85					90					95		
CTC	GTA	GCC	TAC	GCC	GTA	ACA	CTA	AAT	GCG	GGA	GAG	AAG	ATA	ATC	CTC	336

Leu	Val	Ala	Tyr	Ala	Val	Thr	Leu	Asn	Ala	Gly	Glu	Lys	Ile	Ile	Leu	
		100						105					110			
CCA	GAC	CCC	TCT	TAC	CCC	TGT	TAC	AAA	AAC	TTT	GCC	TAC	CTC	TTA	GAC	384
Pro	Asp	Pro	Ser	Tyr	Pro	Cys	Tyr	Lys	Asn	Phe	Ala	Tyr	Leu	Leu	Asp	
		115					120				125					
GCT	CAG	CCG	GTT	TTC	GTA	AAC	GTT	GAC	AAG	GAA	ACG	AAT	TAC	GAA	GTA	432
Ala	Gln	Pro	Val	Phe	Val	Asn	Val	Asp	Lys	Glu	Thr	Asn	Tyr	Glu	Val	
	130					135					140					
AGG	AAA	GAG	ATG	ATA	GAA	GAC	ATT	GAT	GCG	AAA	GCC	CTT	CAC	ATT	TCC	480
Arg	Lys	Glu	Met	Ile	Glu	Asp	Ile	Asp	Ala	Lys	Ala	Leu	His	Ile	Ser	
145					150					155					160	
TCG	CCT	CAA	AAC	CCT	ACG	GGC	ACA	CTC	TAC	TCA	CCT	GAA	ACC	CTG	AAG	528
Ser	Pro	Gln	Asn		Thr	Gly	Thr	Leu	Tyr	Ser	Pro	Glu	Thr		Lys	
				165					170					175		
GAA	CTT	GCG	GAG	TAC	TGC	GAA	GAG	AAG	GGT	ATG	TAC	TTC	ATA	TCC	GAC	576
Glu	Leu	Ala	Glu	Tyr	Cys	Glu	Glu	Lys	Gly	Met	Tyr	Phe	Ile	Ser	Asp	
		180						185					190			
GAG	ATT	TAC	CAC	GGA	CTC	GTT	TAC	GAA	GGT	AGG	GAG	CAC	ACA	GCA	CTT	624
Glu	Ile		Tyr	His	Gly	Leu	Val	Tyr	Glu	Gly	Arg	Glu	His	Thr	Ala	Leu
		195					200					205				
GAG	TTC	TCT	GAC	AGG	GCT	ATT	GTC	ATA	AAC	GGG	TTT	TCT	AAG	TAC	TTC	672
Glu	Phe	Ser	Asp	Arg	Ala	Ile	Val	Ile	Asn	Gly	Phe	Ser	Lys	Tyr	Phe	
	210					215					220					
TGT	ATG	CCA	GGT	TTC	AGG	ATA	GGG	TGG	ATG	ATA	GTT	CCG	GAA	GAA	CTC	720
Cys	Met	Pro	Gly	Phe	Arg	Ile	Gly	Trp	Met	Ile	Val	Pro	Glu	Glu	Leu	
225					230					235					240	
GTG	AGA	AAG	GCG	GAA	ATA	GTA	ATT	CAG	AAC	GTA	TTT	ATA	TCT	GCC	CCG	768
Val	Arg	Lys	Ala	Glu	Ile	Val	Ile	Gln	Asn	Val	Phe	Ile	Ser	Ala	Pro	
			245						250					255		
ACG	CTC	AGT	CAG	TAC	GCC	GCC	CTT	GAG	GCT	TTT	GAT	TAC	GAG	TAT	TTG	816
Thr	Leu	Ser	Gln	Tyr	Ala	Ala	Leu	Glu	Ala	Phe	Asp	Tyr	Glu	Tyr	Leu	
			260					265					270			
GAG	AAG	GTA	AGA	AAA	ACC	TTT	GAA	GAG	AGG	AGG	AAC	TTC	CTT	TAT	GGG	864
Glu	Lys	Val	Arg	Lys	Thr	Phe	Glu	Glu	Arg	Arg	Asn	Phe	Leu	Tyr	Gly	
		275					280					285				
GAA	CTG	AAA	AAA	CTC	TTC	AAG	ATA	GAC	GCG	AAA	CCT	CAG	GGA	GCT	TTT	912
Glu	Leu	Lys	Lys	Leu	Phe	Lys	Ile	Asp	Ala	Lys	Pro	Gln	Gly	Ala	Phe	
	290					295					300					
TAC	GTA	TGG	GCA	AAC	ATA	AGT	GAT	TAC	TCC	ACA	GAT	AGC	TAC	GAA	TTT	960
Tyr	Val	Trp	Ala	Asn	Ile	Ser	Asp	Tyr	Ser	Thr	Asp	Ser	Tyr	Glu	Phe	
305					310					315					320	
GCT	TTA	AAA	CTT	TTA	AGG	GAG	GCG	AGG	GTG	GCG	GTA	ACG	CCC	GGG	GTG	1008
Ala	Leu	Lys	Leu	Leu	Arg	Glu	Ala	Arg	Val	Ala	Val	Thr	Pro	Gly	Val	
				325					330					335		
GAC	TTT	GGA	AAA	AAC	AAA	ACG	AAG	GAG	TAT	ATA	AGG	TTT	GCT	TAT	ACG	1056
Asp	Phe	Gly	Lys	Asn	Lys	Thr	Lys	Glu	Tyr	Ile	Arg	Phe	Ala	Tyr	Thr	
			340					345					350			

AGA AAG ATA GAA GAA CTT AAG GAG GGC GTT GAA AGG ATA AAG AAG TTC	1104
Arg Lys Ile Glu Glu Leu Lys Glu Gly Val Glu Arg Ile Lys Lys Phe	
355 360 365	

TTA GAG AAG CTT AGC TGA	1122
Leu Glu Lys Leu Ser End	
370	

(2) INFORMATION FOR SEQ ID NO:19:

- (i) SEQUENCE CHARACTERISTICS
 (A) LENGTH: 1359 NUCLEOTIDES
 (B) TYPE: NUCLEIC ACID
 (C) STRANDEDNESS: SINGLE
 (D) TOPOLOGY: LINEAR

(ii) MOLECULE TYPE: GENOMIC DNA

(xi) SEQUENCE DESCRIPTION: SEQ ID NO:19:

ATG TGG GAA TTA GAC CCT AAA ACG CTC GAA AAG TGG GAC AAG GAG TAC	48
Met Trp Glu Leu Asp Pro Lys Thr Leu Glu Lys Trp Asp Lys Glu Tyr	
5 10 15	
TTC TGG CAT CCA TTT ACC CAG ATG AAA GTC TAC AGA GAA GAA GAA AAC	96
Phe Trp His Pro Phe Thr Gln Met Lys Val Tyr Arg Glu Glu Glu Asn	
20 25 30	
CTG ATA TTT GAA CGC GGA GAA GGC GTT TAC CTG TGG GAC ATA TAC GGC	144
Leu Ile Phe Glu Arg Gly Glu Gly Val Tyr Leu Trp Asp Ile Tyr Gly	
35 40 45	
AGG AAG TAT ATA GAT GCC ATA TCT TCC CTC TGG TGC AAC GTC CAC GGA	192
Arg Lys Tyr Ile Asp Ala Ile Ser Ser Leu Trp Cys Asn Val His Gly	
50 55 60	
CAT AAC CAC CCT AAA CTG AAC AAC GCA GTT ATG AAA CAG CTC TGT AAG	240
His Asn His Pro Lys Leu Asn Asn Ala Val Met Lys Gln Leu Cys Lys	
65 70 75 80	
GTA GCT CAC ACA ACT ACT CTG GGA AGT TCC AAC GTT CCC GCC ATA CTC	288
Val Ala His Thr Thr Leu Gly Ser Ser Asn Val Pro Ala Ile Leu	
85 90 95	
CTT GCA AAG AAG CTT GTA GAA ATT TCT CCT GAA GGA TTA AAC AAG GTC	336
Leu Ala Lys Lys Leu Val Glu Ile Ser Pro Glu Gly Leu Asn Lys Val	
100 105 110	
TTT TAC TCC GAA GAC GGT GCG GAA GCA GTA GAG ATA GCG ATA AAG ATG	384
Phe Tyr Ser Glu Asp Gly Ala Glu Ala Val Glu Ile Ala Ile Lys Met	
115 120 125	
GCT TAT CAC TAC TGG AAG AAC AAG GGA GTT AAA GGG AAA AAC GTT TTC	432
Ala Tyr His Tyr Trp Lys Asn Lys Gly Val Lys Gly Lys Asn Val Phe	
130 135 140	
ATA ACG CTT TCC GAA GCC TAC CAC GGG GAT ACT GTA GGA GCG GTT AGC	480
Ile Thr Leu Ser Glu Ala Tyr His Gly Asp Thr Val Gly Ala Val Ser	
145 150 155 160	
GTA GGG GGT ATA GAA CTC TTC CAC GGA ACT TAT AAA GAT CTC CTT TTC	528
Val Gly Gly Ile Glu Leu Phe His Gly Thr Tyr Lys Asp Leu Leu Phe	
165 170 175	

AAG Lys	ACT Thr	ATA Ile	AAA Lys 180	CTC Leu	CCA Pro	TCT Ser	CCT Pro	TAC Tyr 185	CTG Leu	TAC Tyr	TGC Cys	AAG Lys	GAA Glu 190	AAG Lys	TAC Tyr	576
GGG Gly	GAA Glu	CTC Leu 195	TGC Cys	CCT Pro	GAG Glu	TGC Cys	ACG Thr 200	GCA Ala	GAT Asp	TTA Leu	TTA Leu	AAA Lys 205	CAA Gln	CTG Leu	GAA Glu	624
GAT Asp	ATC Ile 210	CTG Leu	AAG Lys	TCG Ser	CGG Arg	GAA Glu 215	GAT Asp	ATC Ile	GTT Val	GCG Ala	GTC Val 220	ATT Ile	ATG Met	GAA Glu	GCG Ala	672
GGA Gly 225	ATT Ile	CAG Gln	GCA Ala	GCC Ala	GCG Ala 230	GGA Gly	ATG Met	CTC Leu	CCC Pro	TTC Phe 235	CCT Pro	CCG Pro	GGA Gly	TTT Phe	TTG Leu 240	720
AAA Lys	GGC Gly	GTA Val	AGG Arg	GAG Glu 245	CTT Leu	ACG Thr	AAG Lys	AAA Lys	TAC Tyr 250	GAC Asp	ACT Thr	TTA Leu	ATG Met	ATA Ile 255	GTT Val	768
GAC Asp	GAG Glu	GTT Val	GCC Ala 260	ACG Thr	GGA Gly	TTT Phe	GGC Gly	AGG Arg 265	ACG Thr	GGA Gly	ACG Thr	ATG Met	TTT Phe 270	TAC Tyr	TGT Cys	816
GAG Glu	CAG Gln	GAA Glu 275	GGA Gly	GTC Val	AGT Ser	CCG Pro	GAC Asp 280	TTT Phe	ATG Met	TGT Cys	CTA Leu	GGT Gly 285	AAG Lys	GGT Gly	ATA Ile	864
ACC Thr	GGA Gly 290	GGG Gly	TAC Tyr	CTC Leu	CCG Pro	CTT Leu 295	GCT Ala	GCG Ala	ACA Thr	CTC Leu	ACA Thr 300	ACG Thr	GAC Asp	GAG Glu	GTG Val	912
TTC Phe 305	AAT Asn	GCC Ala	TTT Phe	TTA Leu	GGT Gly 310	GAG Glu	TTC Phe	GGG Gly	GAG Glu	GCA Ala 315	AAG Lys	CAC His	TTT Phe	TAC Tyr	CAC His 320	960
GGG Gly	CAC His	ACC Thr	TAC Tyr	ACT Thr 325	GGA Gly	AAT Asn	AAC Asn	CTC Leu	GCC Ala 330	TGT Cys	TCC Ser	GTT Val	GCA Ala 335	CTC Leu	GCA Ala	1008
AAC Asn	TTA Leu	GAA Glu 340	GTT Val	TTT Phe	GAG Glu	GAA Glu	GAA Glu	AGA Arg 345	ACT Thr	TTA Leu	GAG Glu	AAG Lys	CTC Leu 350	CAA Gln	CCA Pro	1056
AAG Lys	ATA Ile	AAG Lys 355	CTT Leu	TTA Leu	AAG Lys	GAA Glu	AGG Arg 360	CTT Leu	CAG Gln	GAG Glu	TTC Phe	TGG Trp 365	GAA Glu	CTC Leu	AAG Lys	1104
CAC His	GTT Val 370	GGA Gly	GAT Asp	GTT Val	AGA Arg	CAG Gln 375	CTA Leu	GGT Gly	TTT Phe	ATG Met	GCT Ala 380	GGA Gly	ATA Ile	GAG Glu	CTG Leu	1152
GTG Val 385	AAG Lys	GAC Asp	AAA Lys	GAA Glu	AAG Lys 390	GGA Gly	GAA Glu	CCT Pro	TTC Phe	CCT Pro 395	TAC Tyr	GGT Gly	GAA Glu	AGG Arg	ACG Thr 400	1200
GGA Gly	TTT Phe	AAG Lys	GTG Val	GCT Ala 405	TAC Tyr	AAG Lys	TGC Cys	AGG Arg	GAA Glu 410	AAA Lys	GGG Gly	GTG Val	TTT Phe	TTG Leu 415	AGA Arg	1245
CCG Pro	CTC Leu	GGA Gly	GAC Asp	GTT Val	ATG Met	GTA Val	TTG Leu	ATG Met	ATG Met	CCT Pro	CTT Leu	GTA Val	ATA Ile	GAG Glu	GAA Glu	1293

Asn Asp Ile Asp Ser Val Tyr Lys Leu Leu Asp Glu Glu Thr Ala Gly	
165 170 175	
ATA ATT ATT GAA GTT ATA CAA GGA GAG GGC GGA GTA AAC GAG GCG AGT	576
Ile Ile Ile Glu Val Ile Gln Gly Glu Gly Gly Val Asn Glu Ala Ser	
180 185 190	
GAG GAT TTT CTA AGT AAA CTC CAG GAA ATT TGT AAA GAA AAA GAT GTG	624
Glu Asp Phe Leu Ser Lys Leu Gln Glu Ile Cys Lys Glu Lys Asp Val	
195 200 205	
CTC TTA ATT ATA GAC GAA GTG CAA ACG GGA ATA GGA AGG ACC GGG GAA	672
Leu Leu Ile Ile Asp Glu Val Gln Thr Gly Ile Gly Arg Thr Gly Glu	
210 215 220	
TTC TAC GCA TAT CAA CAC TTC AAT CTA AAA CCG GAC GTA ATT GCG CTT	720
Phe Tyr Ala Tyr Gln His Phe Asn Leu Lys Pro Asp Val Ile Ala Leu	
225 230 235 240	
GCG AAG GGA CTC GGA GGA GGT GTG CCA ATA GGT GCC ATC CTT GCA AGG	768
Ala Lys Gly Leu Gly Gly Gly Val Pro Ile Gly Ala Ile Leu Ala Arg	
245 250 255	
GAA GAA GTG GCC CAG AGC TTT ACT CCC GGC TCC CAC GGC TCT ACC TTC	816
Glu Glu Val Ala Gln Ser Phe Thr Pro Gly Ser His Gly Ser Thr Phe	
260 265 270	
GGA GGA AAC CCC TTA GCC TGC AGG GCG GGA ACA GTG GTA GTA GAT GAA	864
Gly Gly Asn Pro Leu Ala Cys Arg Ala Gly Thr Val Val Val Asp Glu	
275 280 285	
GTT GAA AAA CTC CTG CCT CAC GTA AGG GAA GTG GGG AAT TAC TTC AAA	912
Val Glu Lys Leu Leu Pro His Val Arg Glu Val Gly Asn Tyr Phe Lys	
290 295 300	
GAA AAA CTG AAG GAA CTC GGC AAA GGA AAG GTA AAG GGA AGA GGA TTG	960
Glu Lys Leu Lys Glu Leu Gly Lys Gly Lys Val Lys Gly Arg Gly Leu	
305 310 315 320	
ATG CTC GGT CTT GAA CTT GAA AGA GAG TGT AAA GAT TAC GTT CTC AAG	1008
Met Leu Gly Leu Glu Leu Glu Arg Glu Cys Lys Asp Tyr Val Leu Lys	
325 330 335	
GCT CTT GAA AGG GAC TTC TCA TAA	1032
Ala Leu Glu Arg Asp Phe Ser End	
340	

(2) INFORMATION FOR SEQ ID NO:21:

- (i) SEQUENCE CHARACTERISTICS
 (A) LENGTH: 1197 NUCLEOTIDES
 (B) TYPE: NUCLEIC ACID
 (C) STRANDEDNESS: SINGLE
 (D) TOPOLOGY: LINEAR

(ii) MOLECULE TYPE: GENOMIC DNA

(xi) SEQUENCE DESCRIPTION: SEQ ID NO:21:

ATG CGG AAA CTG GCC GAG CGG GCG CAG AAA CTG AGC CCC TCT CCC ACC	48
Met Arg Lys Leu Ala Glu Arg Ala Gln Lys Leu Ser Pro Ser Pro Thr	
5 10 15	

CTC	TCG	GTG	GAC	ACC	AAG	GCC	AAG	GAG	CTT	TTG	CGG	CAG	GGG	GAA	AGG	96
Leu	Ser	Val	Asp	Thr	Lys	Ala	Lys	Glu	Leu	Leu	Arg	Gln	Gly	Glu	Arg	
		20						25					30			
GTC	ATC	AAT	TTC	GGG	GCG	GGG	GAG	CCG	GAC	TTC	GAT	ACA	CCG	GAA	CAC	144
Val	Ile	Asn	Phe	Gly	Ala	Gly	Glu	Pro	Asp	Phe	Asp	Thr	Pro	Glu	His	
		35					40					45				
ATC	AAG	GAA	GCG	GCG	AAG	CGA	GCT	TTA	GAT	CAG	GGC	TTC	ACC	AAG	TAC	192
Ile	Lys	Glu	Ala	Ala	Lys	Arg	Ala	Leu	Asp	Gln	Gly	Phe	Thr	Lys	Tyr	
	50					55					60					
ACG	CCG	GTG	GCT	GGG	ATC	TTA	CCT	CTT	CGG	GAG	GCC	ATA	TGC	GAG	AAG	240
Thr	Pro	Val	Ala	Gly	Ile	Leu	Pro	Leu	Arg	Glu	Ala	Ile	Cys	Glu	Lys	
	65				70				75						80	
CTT	TAC	CGC	GAC	AAT	CAA	CTG	GAA	TAC	AGC	CCG	AAT	GAG	ATC	GTG	GTC	288
Leu	Tyr	Arg	Asp	Asn	Gln	Leu	Glu	Tyr	Ser	Pro	Asn	Glu	Ile	Val	Val	
				85					90					95		
TCC	TGT	GGC	GCC	AAG	CAT	TCT	ATT	TTC	AAC	GCT	CTG	CAG	GTC	CTC	CTG	336
Ser	Cys	Gly	Ala	Lys	His	Ser	Ile	Phe	Asn	Ala	Leu	Gln	Val	Leu	Leu	
			100					105					110			
GAC	CCG	GGG	GAC	GAG	GTG	ATA	ATC	CCC	GTC	CCC	TAC	TGG	ACT	TCC	TAT	384
Asp	Pro	Gly	Asp	Glu	Val	Ile	Ile	Pro	Val	Pro	Tyr	Trp	Thr	Ser	Tyr	
		115					120					125				
CCG	GAG	CAG	GTG	AAG	CTG	GCG	GGA	GGG	GTG	CCG	GTT	TTC	GTC	CCC	ACC	432
Pro	Glu	Gln	Val	Lys	Leu	Ala	Gly	Gly	Val	Pro	Val	Phe	Val	Pro	Thr	
	130					135					140					
TCT	CCC	GAG	AAC	GAC	TTC	AAG	CTC	AGG	CCG	GAA	GAT	CTA	CGT	GCG	GCT	480
Ser	Pro	Glu	Asn	Asp	Phe	Lys	Leu	Arg	Pro	Glu	Asp	Leu	Arg	Ala	Ala	
	145				150					155					160	
GTA	ACC	CCG	CGC	ACC	CGC	CTT	TTG	ATC	CTC	AAT	TCC	CCG	GCC	AAC	CCC	528
Val	Thr	Pro	Arg	Thr	Arg	Leu	Leu	Ile	Leu	Asn	Ser	Pro	Ala	Asn	Pro	
				165					170					175		
ACA	GGC	ACC	GTT	TAC	CGC	CGG	GAG	GAA	CTT	ATC	GGC	TTA	GCG	GAG	GTA	576
Thr	Gly	Thr	Val	Tyr	Arg	Arg	Glu	Glu	Leu	Ile	Gly	Leu	Ala	Glu	Val	
			180				185					190				
GCC	CTG	GAG	GCC	GAC	CTA	TGG	ATC	TTG	TCG	GAC	GAG	ATC	TAC	GAA	AAG	624
Ala	Leu	Glu	Ala	Asp	Leu	Trp	Ile	Leu	Ser	Asp	Glu	Ile	Tyr	Glu	Lys	
		195					200					205				
CTG	ATC	TAC	GAC	GGG	ATG	GAG	CAC	GTG	AGC	ATA	GCC	GCG	CTC	GAC	CCG	672
Leu	Ile	Tyr	Asp	Gly	Met	Glu	His	Val	Ser	Ile	Ala	Ala	Leu	Asp	Pro	
	210					215					220					
GAG	GTC	AAA	AAG	CGC	ACG	ATT	GTG	GTA	AAC	GGT	GTT	TCC	AAG	GCT	TAC	720
Glu	Val	Lys	Lys	Arg	Thr	Ile	Val	Val	Asn	Gly	Val	Ser	Lys	Ala	Tyr	
	225				230					235					240	
GCC	ATG	ACC	GGT	TGG	CGC	ATA	GGT	TAT	GCT	GCC	GCT	CCC	CGG	CCG	ATA	768
Ala	Met	Thr	Gly	Trp	Arg	Ile	Gly	Tyr	Ala	Ala	Ala	Pro	Arg	Pro	Ile	
				245				250						255		
GCC	CAG	GCC	ATG	ACC	AAC	CTC	CAA	AGC	CAC	AGT	ACC	TCT	AAC	CCC	ACT	816
Ala	Gln	Ala	Met	Thr	Asn	Leu	Gln	Ser	His	Ser	Thr	Ser	Asn	Pro	Thr	

260	265	270	
TCC GTA GCC CAG GCG GCG GCG CTG GCC GCT CTG AAG GGG CCA CAA GAG Ser Val Ala Gln Ala Ala Ala Leu Ala Ala Leu Lys Gly Pro Gln Glu 275 280 285			864
CCG GTG GAG AAC ATG CGC CGG GCT TTT CAA AAG CGG CGG GAT TTC ATC Pro Val Glu Asn Met Arg Arg Ala Phe Gln Lys Arg Arg Asp Phe Ile 290 295 300			912
TGG CAG TAC CTA AAC TCC TTA CCC GGA GTG CGC TGC CCC AAA CCT TTA Trp Gln Tyr Leu Asn Ser Leu Pro Gly Val Arg Cys Pro Lys Pro Leu 305 310 315 320			960
GGG GCC TTT TAC GTC TTT CCA GAA GTT GAG CGG GCT TTT GGG CCG CCG Gly Ala Phe Tyr Val Phe Pro Glu Val Glu Arg Ala Phe Gly Pro Pro 325 330 335			1008
TCT AAA AGG ACG GGA AAT ACT ACC GCT AGC GAC CTG GCC CTT TTC CTC Ser Lys Arg Thr Gly Asn Thr Thr Ala Ser Asp Leu Ala Leu Phe Leu 340 345 350			1056
CTG GAA GAG ATA AAA GTG GCC ACC GTG GCT GGG GCT GCC TTT GGG GAC Leu Glu Glu Ile Lys Val Ala Thr Val Ala Gly Ala Ala Phe Gly Asp 355 360 365			1104
GAT CGC TAC CTG CGC TTT TCC TAC GCC CTG CGG CTG GAA GAT ATC GAA Asp Arg Tyr Leu Arg Phe Ser Tyr Ala Leu Arg Leu Glu Asp Ile Glu 370 375 380			1152
GAG GGG ATG CAA CGG TTT AAA GAA TTG ATC GAA GCG GCA CTT TAA Glu Gly Met Gln Arg Phe Lys Glu Leu Ile Glu Ala Ala Leu End 385 390 395			1197

- (2) INFORMATION FOR SEQ ID NO:22:
- (i) SEQUENCE CHARACTERISTICS
- (A) LENGTH: 1779 NUCLEOTIDES
- (B) TYPE: NUCLEIC ACID
- (C) STRANDEDNESS: SINGLE
- (D) TOPOLOGY: LINEAR
- (ii) MOLECULE TYPE: GENOMIC DNA
- (xi) SEQUENCE DESCRIPTION: SEQ ID NO:22:

ATG TGC GGG ATA GTC GGA TAC GTA GGG AGG GAT TTA GCC CTT CCT ATA Met Cys Gly Ile Val Gly Tyr Val Gly Arg Asp Leu Ala Leu Pro Ile 5 10 15	48
GTC CTC GGA GCT CTT GAG AGA CTC GAA TAC AGG GGT TAC GAC TCC GCG Val Leu Gly Ala Leu Glu Arg Leu Glu Tyr Arg Gly Tyr Asp Ser Ala 20 25 30	96
GGA GTT GCC CTT ATA GAA GAC GGG AAA CTC ATA GTT GAA AAG AAG AAG Gly Val Ala Leu Ile Glu Asp Gly Lys Leu Ile Val Glu Lys Lys Lys 35 40 45	144
GGA AAG ATA AGG GAA CTC GTT AAA GCG CTA TGG GGA AAG GAT TAC AAG Gly Lys Ile Arg Glu Leu Val Lys Ala Leu Trp Gly Lys Asp Tyr Lys 50 55 60	192

GCT AAA ACG GGT ATA GGT CAC ACA CGC TGG GCA ACC CAC GGA AAG CCC Ala Lys Thr Gly Ile Gly His Thr Arg Trp Ala Thr His Gly Lys Pro 65 70 75 80	240
ACG GAC GAG AAC GCC CAC CCC CAC ACC GAC GAA AAA GGT GAG TTT GCA Thr Asp Glu Asn Ala His Pro His Thr Asp Glu Lys Gly Glu Phe Ala 85 90 95	288
GTA GTT CAC AAC GGG ATA ATA GAA AAC TAC TTA GAA CTA AAA GAG GAA Val Val His Asn Gly Ile Ile Glu Asn Tyr Leu Glu Leu Lys Glu Glu 100 105 110	336
CTA AAG AAG GAA GGT GTA AAG TTC AGG TCC GAA ACA GAC ACA GAA GTT Leu Lys Lys Glu Gly Val Lys Phe Arg Ser Glu Thr Asp Thr Glu Val 115 120 125	384
ATA GCC CAC CTC ATA GCG AAG AAC TAC AGG GGG GAC TTA CTG GAG GCC Ile Ala His Leu Ile Ala Lys Asn Tyr Arg Gly Asp Leu Leu Glu Ala 130 135 140	432
GTT TTA AAA ACC GTA AAG AAA TTA AAG GGT GCT TTT GCC TTT GCG GTT Val Leu Lys Thr Val Lys Lys Leu Lys Gly Ala Phe Ala Phe Ala Val 145 150 155 160	480
ATA ACG GTT CAC GAA CCA AAC AGA CTA ATA GGA GTG AAG CAG GGG AGT Ile Thr Val His Glu Pro Asn Arg Leu Ile Gly Val Lys Gln Gly Ser 165 170 175	528
CCT TTA ATC GTC GGA CTC GGA GAA GGA GAA AAC TTC CTC GCT TCA GAT Pro Leu Ile Val Gly Leu Gly Glu Gly Glu Asn Phe Leu Ala Ser Asp 180 185 190	576
ATT CCC GCA ATA CTT CCT TAC ACG AAA AAG ATT ATT GTT CTT GAT GAC Ile Pro Ala Ile Leu Pro Tyr Thr Lys Lys Ile Ile Val Leu Asp Asp 195 200 205	624
GGG GAA ATA GCG GAC CTG ACT CCC GAC ACT GTG AAC ATT TAC AAC TTT Gly Glu Ile Ala Asp Leu Thr Pro Asp Thr Val Asn Ile Tyr Asn Phe 210 215 220	672
GAG GGA GAG CCC GTT TCA AAG GAA GTA ATG ATT ACG CCC TGG GAT CTT Glu Gly Glu Pro Val Ser Lys Glu Val Met Ile Thr Pro Trp Asp Leu 225 230 235 240	720
GTT TCT GCG GAA AAG GGT GGT TTT AAA CAC TTC ATG CTA AAA GAG ATA Val Ser Ala Glu Lys Gly Gly Phe Lys His Phe Met Leu Lys Glu Ile 245 250 255	768
TAC GAA CAG CCC AAA GCC ATA AAC GAC ACA CTC AAG GGT TTC CTC TCA Tyr Glu Gln Pro Lys Ala Ile Asn Asp Thr Leu Lys Gly Phe Leu Ser 260 265 270	816
ACC GAA GAC GCA ATA CCC TTT AAG TTA AAA GAC TTC AGA AGG GTT TTA Thr Glu Asp Ala Ile Pro Phe Lys Leu Lys Asp Phe Arg Arg Val Leu 275 280 285	864
ATA ATA GCG TGC GGG ACC TCT TAC CAC GCG GGC TTC GTC GGA AAG TAC Ile Ile Ala Cys Gly Thr Ser Tyr His Ala Gly Phe Val Gly Lys Tyr 290 295 300	912
TGG ATA GAG AGA TTT GCA GGT GTT CCC ACA GAG GTA ATT TAC GCT TCG Trp Ile Glu Arg Phe Ala Gly Val Pro Thr Glu Val Ile Tyr Ala Ser	960

305	310	315	320	
GAA TTC AGG TAT GCG GAC GTT CCC GTT TCG GAC AAG GAT ATC GTT ATC Glu Phe Arg Tyr Ala Asp Val Pro Val Ser Asp Lys Asp Ile Val Ile 325 330 335				1008
GGA ATT TCC CAG TCA GGA GAG ACC GCT GAC ACA AAG TTT GCC CTT CAG Gly Ile Ser Gln Ser Gly Glu Thr Ala Asp Thr Lys Phe Ala Leu Gln 340 345 350				1056
TCC GCA AAG GAA AAG GGA GCC TTT ACC GTG GGA CTC GTA AAC GTA GTG Ser Ala Lys Glu Lys Gly Ala Phe Thr Val Gly Leu Val Asn Val Val 355 360 365				1104
GGA AGT GCC ATA GAC AGG GAG TCG GAC TTT TCC CTT CAC ACA CAT GCG Gly Ser Ala Ile Asp Arg Glu Ser Asp Phe Ser Leu His Thr His Ala 370 375 380				1152
GGA CCC GAA ATA GGC GTG GCG GCT ACA AAG ACC TTC ACC GCA CAG TTC Gly Pro Glu Ile Gly Val Ala Ala Thr Lys Thr Phe Thr Ala Gln Phe 385 390 395 400				1200
ACC GCA CTC TAC GCC CTT TCG GTA AGG GAA AGT GAG GAG AGG GAA AAT Thr Ala Leu Tyr Ala Leu Ser Val Arg Glu Ser Glu Glu Arg Glu Asn 405 410 415				1248
CTA ATA AGA CTC CTT GAA AAG GTT CCA TCA CTC GTT GAA CAA ACA CTG Leu Ile Arg Leu Leu Glu Lys Val Pro Ser Leu Val Glu Gln Thr Leu 420 425 430				1296
AAC ACC GCA GAA GAA GTG GAG AAG GTA GCG GAA AAG TAC ATG AAA AAG Asn Thr Ala Glu Glu Val Glu Lys Val Ala Glu Lys Tyr Met Lys Lys 435 440 445				1344
AAA AAC ATG CTT TAC CTC GGA AGG TAC TTA AAT TAC CCC ATA GCG CTG Lys Asn Met Leu Tyr Leu Gly Arg Tyr Leu Asn Tyr Pro Ile Ala Leu 450 455 460				1392
GAG GGA GCT CTT AAA CTT AAA GAA ATT TCT TAC ATA CAC GCG GAA GGT Glu Gly Ala Leu Lys Leu Lys Glu Ile Ser Tyr Ile His Ala Glu Gly 465 470 475 480				1440
TAT CCC GCA GGG GAG ATG AAG CAC GGT CCC ATA GCC CTC ATA GAC GAA Tyr Pro Ala Gly Glu Met Lys His Gly Pro Ile Ala Leu Ile Asp Glu 485 490 495				1488
AAC ATG CCG GTT GTG GTA ATC GCA CCG AAA GAC AGG GTT TAC GAG AAG Asn Met Pro Val Val Val Ile Ala Pro Lys Asp Arg Val Tyr Glu Lys 500 505 510				1536
ATA CTC TCA AAC GTA GAA GAG GTT CTC GCA AGA AAG GGA AGG GTT ATT Ile Leu Ser Asn Val Glu Glu Val Leu Ala Arg Lys Gly Arg Val Ile 515 520 525				1584
TCT GTA GGC TTT AAA GGA GAC GAA ACT CTC AAA AGC AAA TCC GAG AGC Ser Val Gly Phe Lys Gly Asp Glu Thr Leu Lys Ser Lys Ser Glu Ser 530 535 540				1632
GTT ATG GAA ATC CCG AAG GCA GAA GAA CCG ATA ACT CCT TTC TTG ACG Val Met Glu Ile Pro Lys Ala Glu Glu Pro Ile Thr Pro Phe Leu Thr 545 550 555 560				1680
GTA ATA CCC CTG CAA CTC TTT GCC TAC TTT ATA GCG AGC AAA CTG GGA Val Ile Pro Leu Gln Leu Phe Ala Tyr Phe Ile Ala Ser Lys Leu Gly				1728

	565	570	575	580
CTG GAT GTG GAT CAG CCG AGA AAT CTC GCC AAA ACG GTC ACG GTG GAA				1776
Leu Asp Val Asp Gln Pro Arg Asn Leu Ala Lys Thr Val Thr Val Glu				
	580	585	590	
TAA				1779
End				

(2) INFORMATION FOR SEQ ID NO:23:

- (i) SEQUENCE CHARACTERISTICS
 (A) LENGTH: 1065 NUCLEOTIDES
 (B) TYPE: NUCLEIC ACID
 (C) STRANDEDNESS: SINGLE
 (D) TOPOLOGY: LINEAR

(ii) MOLECULE TYPE: GENOMIC DNA

(xi) SEQUENCE DESCRIPTION: SEQ ID NO:23:

ATG ATA CCC CAG AGG ATT AAG GAA CTT GAA GCT TAC AAG ACG GAG GTC	48
Met Ile Pro Gln Arg Ile Lys Glu Leu Glu Ala Tyr Lys Thr Glu Val	
	5 10 15
ACT CCC GCC TCC GTC AGG CTT TCC TCT AAC GAA TTC CCC TAC GAC TTT	96
Thr Pro Ala Ser Val Arg Leu Ser Ser Asn Glu Phe Pro Tyr Asp Phe	
	20 25 30
CCC GAG GAG ATA AAA CAA AGG GCC TTA GAA GAA TTA AAA AAG GTT CCC	144
Pro Glu Glu Ile Lys Gln Arg Ala Leu Glu Glu Leu Lys Lys Val Pro	
	35 40 45
TTG AAC AAA TAC CCA GAC CCC GAA GCG AAA GAG TTA AAA GCG GTT CTT	192
Leu Asn Lys Tyr Pro Asp Pro Glu Ala Lys Glu Leu Lys Ala Val Leu	
	50 55 60
GCG GAT TTT TTC GGC GTT AAG GAA GAA AAT TTA GTT CTC GGT AAC GGT	240
Ala Asp Phe Phe Gly Val Lys Glu Glu Asn Leu Val Leu Gly Asn Gly	
	65 70 75 80
TCG GAC GAA CTC ATA TAC TAC CTC TCA ATA GCT ATA GGT GAA CTT TAC	288
Ser Asp Glu Leu Ile Tyr Tyr Leu Ser Ile Ala Ile Gly Glu Leu Tyr	
	85 90 95
ATA CCC GTT TAC ATA CCT GTT CCC ACC TTT CCC ATG TAC GAG ATA AGT	336
Ile Pro Val Tyr Ile Pro Val Pro Thr Phe Pro Met Tyr Glu Ile Ser	
	100 105 110
GCG AAA GTT CTC GGA AGA CCC CTC GTA AAG GTT CAA CTG GAC GAA AAC	384
Ala Lys Val Leu Gly Arg Pro Leu Val Lys Val Gln Leu Asp Glu Asn	
	115 120 125
TTT GAT ATA GAC TTA GAA AGA AGT ATT GAA TTA ATA GAG AAA GAA AAA	432
Phe Asp Ile Asp Leu Glu Arg Ser Ile Glu Leu Ile Glu Lys Glu Lys	
	130 135 140
CCC GTT CTC GGG TAC TTT GCT TAC CCA AAC AAC CCC ACG GGA AAC CTC	480
Pro Val Leu Gly Tyr Phe Ala Tyr Pro Asn Asn Pro Thr Gly Asn Leu	
	145 150 155 160
TTT TCC AGG GGA AAG ATT GAG GAG ATA AGA AAC AGG GGT GTT TTC TGT	528
Phe Ser Arg Gly Lys Ile Glu Glu Ile Arg Asn Arg Gly Val Phe Cys	

Met	Lys	Pro	Tyr	Ala	Lys	Tyr	Ile	Trp	Leu	Asp	Gly	Arg	Ile	Leu	Lys		
				5					10					15			
TGG	GAA	GAC	GCG	AAA	ATA	CAC	GTG	TTG	ACT	CAC	GCG	CTT	CAC	TAC	GGA		96
Trp	Glu	Asp	Ala	Lys	Ile	His	Val	Leu	Thr	His	Ala	Leu	His	Tyr	Gly		
			20					25					30				
ACC	TCT	ATA	TTC	GAG	GGA	ATA	AGA	GGG	TAT	TGG	AAC	GGC	GAT	AAT	TTG		144
Thr	Ser	Ile	Phe	Glu	Gly	Ile	Arg	Gly	Tyr	Trp	Asn	Gly	Asp	Asn	Leu		
		35					40					45					
CTC	GTC	TTT	AGG	TTA	GAA	GAA	CAC	ATC	GAC	CGC	ATG	TAC	AGA	TCG	GCT		192
Leu	Val	Phe	Arg	Leu	Glu	Glu	His	Ile	Asp	Arg	Met	Tyr	Arg	Ser	Ala		
		50					55				60						
AAG	ATA	CTA	GGC	ATA	AAT	ATT	CCG	TAT	ACA	AGA	GAG	GAA	GTC	CGC	CAA		240
Lys	Ile	Leu	Gly	Ile	Asn	Ile	Pro	Tyr	Thr	Arg	Glu	Glu	Val	Arg	Gln		
	65				70					75					80		
GCT	GTA	CTA	GAG	ACC	ATA	AAG	GCT	AAT	AAC	TTC	CGA	GAG	GAT	GTC	TAC		288
Ala	Val	Leu	Glu	Thr	Ile	Lys	Ala	Asn	Asn	Phe	Arg	Glu	Asp	Val	Tyr		
				85				90						95			
ATA	AGA	CCT	GTG	GCG	TTT	GTC	GCC	TCG	CAG	ACG	GTG	ACG	CTT	GAC	ATA		336
Ile	Arg	Pro	Val	Ala	Phe	Val	Ala	Ser	Gln	Thr	Val	Thr	Leu	Asp	Ile		
			100					105					110				
AGA	AAT	TTG	GAA	GTC	TCC	CTC	GCG	GTT	ATT	GTA	TTC	CCA	TTT	GGC	AAA		384
Arg	Asn	Leu	Glu	Val	Ser	Leu	Ala	Val	Ile	Val	Phe	Pro	Phe	Gly	Lys		
		115					120					125					
TAC	CTC	TCG	CCC	AAC	GGC	ATT	AAG	GCA	ACG	ATT	GTA	AGC	TGG	CGT	AGA		432
Tyr	Leu	Ser	Pro	Asn	Gly	Ile	Lys	Ala	Thr	Ile	Val	Ser	Trp	Arg	Arg		
		130				135					140						
GTA	CAT	AAT	ACA	ATG	CTC	CCT	GTG	ATG	GCA	AAA	ATC	GGC	GGT	ATA	TAT		480
Val	His	Asn	Thr	Met	Leu	Pro	Val	Met	Ala	Lys	Ile	Gly	Gly	Ile	Tyr		
	145				150					155					160		
GTA	AAC	TCT	GTA	CTT	GCG	CTT	GTA	GAG	GCT	AGA	AGC	AGG	GGA	TTT	GAC		528
Val	Asn	Ser	Val	Leu	Ala	Leu	Val	Glu	Ala	Arg	Ser	Arg	Gly	Phe	Asp		
			165					170						175			
GAG	GCT	TTA	TTA	ATG	GAC	GTT	AAC	GGT	TAT	GTT	GTT	GAG	GGT	TCT	GGA		576
Glu	Ala	Leu	Leu	Met	Asp	Val	Asn	Gly	Tyr	Val	Val	Glu	Gly	Ser	Gly		
			180					185					190				
GAG	AAT	ATT	TTC	ATT	GTC	AGA	GGT	GGA	AGG	CTT	TTC	ACG	CCG	CCA	GTA		624
Glu	Asn	Ile	Phe	Ile	Val	Arg	Gly	Gly	Arg	Leu	Phe	Thr	Pro	Pro	Val		
		195					200					205					
CAC	GAA	TCT	ATC	CTC	GAG	GGA	ATT	ACG	AGG	GAT	ACG	GTA	ATA	AAG	CTC		672
His	Glu	Ser	Ile	Leu	Glu	Gly	Ile	Thr	Arg	Asp	Thr	Val	Ile	Lys	Leu		
		210				215					220						
AGC	GGG	GAT	GTG	GGA	CTT	CGG	GTG	GAG	GAA	AAG	CCT	ATT	ACG	AGG	GAG		720
Ser	Gly	Asp	Val	Gly	Leu	Arg	Val	Glu	Glu	Lys	Pro	Ile	Thr	Arg	Glu		
	225				230					235					240		
GAG	GTG	TAT	ACA	GCC	GAC	GAG	GTG	TTT	TTA	GTA	GGA	ACC	GCC	GCA	GAG		768
Glu	Val	Tyr	Thr	Ala	Asp	Glu	Val	Phe	Leu	Val	Gly	Thr	Ala	Ala	Glu		
				245					250					255			

ATA ACG CCA GTG GTG GAG GTT GAC GGC AGA ACA ATC GGC ACA GGC AAG	816
Ile Thr Pro Val Val Glu Val Asp Gly Arg Thr Ile Gly Thr Gly Lys	
260 265 270	
CCG GGC CCC ATT ACG ACA AAA ATA GCT GAG CTG TAC TCA AAC GTC GTG	864
Pro Gly Pro Ile Thr Thr Lys Ile Ala Glu Leu Tyr Ser Asn Val Val	
275 280 285	
AGA GGC AAA GTA GAG AAA TAC TTA AAT TGG ATC ACT CCT GTG TAT TAG	912
Arg Gly Lys Val Glu Lys Tyr Leu Asn Trp Ile Thr Pro Val Tyr End	
290 295 300	

- (2) INFORMATION FOR SEQ ID NO:25:
- (i) SEQUENCE CHARACTERISTICS
- (A) LENGTH: 414 AMINO ACIDS
- (B) TYPE: AMINO ACID
- (D) TOPOLOGY: LINEAR
- (ii) MOLECULE TYPE: PROTEIN
- (xi) SEQUENCE DESCRIPTION: SEQ ID NO:25:

Met Ile Glu Asp Pro Met Asp Trp Ala Phe Pro Arg Ile Lys Arg Leu	
5 10 15	
Pro Gln Tyr Val Phe Ser Leu Val Asn Glu Leu Lys Tyr Lys Leu Arg	
20 25 30	
Arg Glu Gly Glu Asp Val Val Asp Leu Gly Met Gly Asn Pro Asn Met	
35 40 45	
Pro Pro Ala Lys His Ile Ile Asp Lys Leu Cys Glu Val Ala Gln Lys	
50 55 60	
Pro Asn Val His Gly Tyr Ser Ala Ser Arg Gly Ile Pro Arg Leu Arg	
65 70 75 80	
Lys Ala Ile Cys Asn Phe Tyr Glu Glu Arg Tyr Gly Val Lys Leu Asp	
85 90 95	
Pro Glu Arg Glu Ala Ile Leu Thr Ile Gly Ala Lys Glu Gly Tyr Ser	
100 105 110	
His Leu Met Leu Ala Met Ile Ser Pro Gly Asp Thr Val Ile Val Pro	
115 120 125	
Asn Pro Thr Tyr Pro Ile His Tyr Tyr Ala Pro Ile Ile Ala Gly Gly	
130 135 140	
Glu Val His Ser Ile Pro Leu Asn Phe Ser Asp Asp Gln Asp His Gln	
145 150 155 160	
Glu Glu Phe Leu Arg Arg Leu Tyr Glu Ile Val Lys Thr Ala Met Pro	
165 170 175	
Lys Pro Lys Ala Val Val Ile Ser Phe Pro His Asn Pro Thr Thr Ile	
180 185 190	
Thr Val Glu Lys Asp Phe Phe Lys Glu Ile Val Lys Phe Ala Lys Glu	
195 200 205	

His Gly Leu Trp Ile Ile His Asp Phe Ala Tyr Ala Asp Ile Ala Phe
 210 215 220
 Asp Gly Tyr Lys Pro Pro Ser Ile Leu Glu Ile Glu Gly Ala Lys Asp
 225 230 235 240
 Val Ala Val Glu Leu Tyr Ser Met Ser Lys Gly Phe Ser Met Ala Gly
 245 250 255
 Trp Arg Val Ala Phe Val Val Gly Asn Glu Ile Leu Ile Lys Asn Leu
 260 265 270
 Ala His Leu Lys Ser Tyr Leu Asp Tyr Gly Ile Phe Thr Pro Ile Gln
 275 280 285
 Val Ala Ser Ile Ile Ala Leu Glu Ser Pro Tyr Glu Ile Val Glu Lys
 290 295 300
 Thr Ala Lys Val Tyr Gln Lys Arg Arg Asp Val Leu Val Glu Gly Leu
 305 310 315 320
 Asn Arg Leu Gly Trp Lys Val Lys Lys Pro Lys Ala Thr Met Phe Val
 325 330 335
 Trp Ala Lys Ile Pro Glu Trp Ile Asn Met Asn Ser Leu Asp Phe Ser
 340 345 350
 Leu Phe Leu Leu Lys Glu Ala Lys Val Ala Val Ser Pro Gly Val Gly
 355 360 365
 Phe Gly Gln Tyr Gly Glu Gly Tyr Val Arg Phe Ala Leu Val Glu Asn
 370 375 380
 Glu His Arg Ile Arg Gln Ala Ile Arg Gly Ile Arg Lys Ala Phe Arg
 385 390 395 400
 Lys Leu Gln Lys Glu Arg Lys Leu Glu Pro Glu Arg Ser Ala End
 405 410 414

(2) INFORMATION FOR SEQ ID NO:26:

- (i) SEQUENCE CHARACTERISTICS
 (A) LENGTH: 373 AMINO ACIDS
 (B) TYPE: AMINO ACID
 (D) TOPOLOGY: LINEAR

(ii) MOLECULE TYPE: PROTEIN

(xi) SEQUENCE DESCRIPTION: SEQ ID NO:26:

Met Asp Arg Leu Glu Lys Val Ser Pro Phe Ile Val Met Asp Ile Leu
 5 10 15
 Ala Gln Ala Gln Lys Tyr Glu Asp Val Val His Met Glu Ile Gly Glu
 20 25 30
 Pro Asp Leu Glu Pro Ser Pro Lys Val Met Glu Ala Leu Glu Arg Ala
 35 40 45
 Val Lys Glu Lys Thr Phe Phe Tyr Thr Pro Ala Leu Gly Leu Trp Glu
 50 55 60
 Leu Arg Glu Arg Ile Ser Glu Phe Tyr Arg Lys Lys Tyr Ser Val Glu

65					70					75				80
Val	Ser	Pro	Glu	Arg	Val	Ile	Val	Thr	Thr	Gly	Thr	Ser	Gly	Ala
				85					90					95
Leu	Val	Ala	Tyr	Ala	Val	Thr	Leu	Asn	Ala	Gly	Glu	Lys	Ile	Ile
			100					105					110	Leu
Pro	Asp	Pro	Ser	Tyr	Pro	Cys	Tyr	Lys	Asn	Phe	Ala	Tyr	Leu	Leu
		115					120					125	Asp	
Ala	Gln	Pro	Val	Phe	Val	Asn	Val	Asp	Lys	Glu	Thr	Asn	Tyr	Glu
	130					135					140			Val
Arg	Lys	Glu	Met	Ile	Glu	Asp	Ile	Asp	Ala	Lys	Ala	Leu	His	Ile
145					150					155				160
Ser	Pro	Gln	Asn	Pro	Thr	Gly	Thr	Leu	Tyr	Ser	Pro	Glu	Thr	Leu
			165						170					175
Glu	Leu	Ala	Glu	Tyr	Cys	Glu	Glu	Lys	Gly	Met	Tyr	Phe	Ile	Ser
			180					185					190	Asp
Glu	Ile	Tyr	His	Gly	Leu	Val	Tyr	Glu	Gly	Arg	Glu	His	Thr	Ala
		195					200					205		Leu
Glu	Phe	Ser	Asp	Arg	Ala	Ile	Val	Ile	Asn	Gly	Phe	Ser	Lys	Tyr
	210					215					220			Phe
Cys	Met	Pro	Gly	Phe	Arg	Ile	Gly	Trp	Met	Ile	Val	Pro	Glu	Glu
225					230					235				Leu
Val	Arg	Lys	Ala	Glu	Ile	Val	Ile	Gln	Asn	Val	Phe	Ile	Ser	Ala
				245					250					255
Thr	Leu	Ser	Gln	Tyr	Ala	Ala	Leu	Glu	Ala	Phe	Asp	Tyr	Glu	Tyr
			260					265					270	Leu
Glu	Lys	Val	Arg	Lys	Thr	Phe	Glu	Glu	Arg	Arg	Asn	Phe	Leu	Tyr
		275					280					285		Gly
Glu	Leu	Lys	Lys	Leu	Phe	Lys	Ile	Asp	Ala	Lys	Pro	Gln	Gly	Ala
	290					295					300			Phe
Tyr	Val	Trp	Ala	Asn	Ile	Ser	Asp	Tyr	Ser	Thr	Asp	Ser	Tyr	Glu
305					310					315				320
Ala	Leu	Lys	Leu	Leu	Arg	Glu	Ala	Arg	Val	Ala	Val	Thr	Pro	Gly
				325					330					335
Asp	Phe	Gly	Lys	Asn	Lys	Thr	Lys	Glu	Tyr	Ile	Arg	Phe	Ala	Tyr
			340					345					350	Thr
Arg	Lys	Ile	Glu	Glu	Leu	Lys	Glu	Gly	Val	Glu	Arg	Ile	Lys	Lys
		355					360					365		Phe
Leu	Glu	Lys	Leu	Ser										
	370													

(2) INFORMATION FOR SEQ ID NO:27:

(i) SEQUENCE CHARACTERISTICS

(A) LENGTH: 453 AMINO ACIDS

(B) TYPE: AMINO ACID

(D) TOPOLOGY: LINEAR

(ii) MOLECULE TYPE: PROTEIN

(xi) SEQUENCE DESCRIPTION: SEQ ID NO:27:

Met Trp Glu Leu Asp Pro Lys Thr Leu Glu Lys Trp Asp Lys Glu Tyr
5 10 15
Phe Trp His Pro Phe Thr Gln Met Lys Val Tyr Arg Glu Glu Glu Asn
20 25 30
Leu Ile Phe Glu Arg Gly Glu Gly Val Tyr Leu Trp Asp Ile Tyr Gly
35 40 45
Arg Lys Tyr Ile Asp Ala Ile Ser Ser Leu Trp Cys Asn Val His Gly
50 55 60
His Asn His Pro Lys Leu Asn Asn Ala Val Met Lys Gln Leu Cys Lys
65 70 75 80
Val Ala His Thr Thr Thr Leu Gly Ser Ser Asn Val Pro Ala Ile Leu
85 90 95
Leu Ala Lys Lys Leu Val Glu Ile Ser Pro Glu Gly Leu Asn Lys Val
100 105 110
Phe Tyr Ser Glu Asp Gly Ala Glu Ala Val Glu Ile Ala Ile Lys Met
115 120 125
Ala Tyr His Tyr Trp Lys Asn Lys Gly Val Lys Gly Lys Asn Val Phe
130 135 140
Ile Thr Leu Ser Glu Ala Tyr His Gly Asp Thr Val Gly Ala Val Ser
145 150 155 160
Val Gly Gly Ile Glu Leu Phe His Gly Thr Tyr Lys Asp Leu Leu Phe
165 170 175
Lys Thr Ile Lys Leu Pro Ser Pro Tyr Leu Tyr Cys Lys Glu Lys Tyr
180 185 190
Gly Glu Leu Cys Pro Glu Cys Thr Ala Asp Leu Leu Lys Gln Leu Glu
195 200 205
Asp Ile Leu Lys Ser Arg Glu Asp Ile Val Ala Val Ile Met Glu Ala
210 215 220
Gly Ile Gln Ala Ala Ala Gly Met Leu Pro Phe Pro Pro Gly Phe Leu
225 230 235 240
Lys Gly Val Arg Glu Leu Thr Lys Lys Tyr Asp Thr Leu Met Ile Val
245 250 255
Asp Glu Val Ala Thr Gly Phe Gly Arg Thr Gly Thr Met Phe Tyr Cys
260 265 270

Glu Gln Glu Gly Val Ser Pro Asp Phe Met Cys Leu Gly Lys Gly Ile
 275 280 285
 Thr Gly Gly Tyr Leu Pro Leu Ala Ala Thr Leu Thr Thr Asp Glu Val
 290 295 300
 Phe Asn Ala Phe Leu Gly Glu Phe Gly Glu Ala Lys His Phe Tyr His
 305 310 315 320
 Gly His Thr Tyr Thr Gly Asn Asn Leu Ala Cys Ser Val Ala Leu Ala
 325 330 335
 Asn Leu Glu Val Phe Glu Glu Glu Arg Thr Leu Glu Lys Leu Gln Pro
 340 345 350
 Lys Ile Lys Leu Leu Lys Glu Arg Leu Gln Glu Phe Trp Glu Leu Lys
 355 360 365
 His Val Gly Asp Val Arg Gln Leu Gly Phe Met Ala Gly Ile Glu Leu
 370 375 380
 Val Lys Asp Lys Glu Lys Gly Glu Pro Phe Pro Tyr Gly Glu Arg Thr
 385 390 395 400
 Gly Phe Lys Val Ala Tyr Lys Cys Arg Glu Lys Gly Val Phe Leu Arg
 405 410 415
 Pro Leu Gly Asp Val Met Val Leu Met Met Pro Leu Val Ile Glu Glu
 420 425 430
 Asp Glu Met Asn Tyr Val Ile Asp Thr Leu Lys Trp Ala Ile Lys Glu
 435 440 445
 Leu Glu Lys Glu Val
 450

- (2) INFORMATION FOR SEQ ID NO:28:
- (i) SEQUENCE CHARACTERISTICS
 - (A) LENGTH: 343 AMINO ACIDS
 - (B) TYPE: AMINO ACID
 - (D) TOPOLOGY: LINEAR
 - (ii) MOLECULE TYPE: PROTEIN
 - (xi) SEQUENCE DESCRIPTION: SEQ ID NO:28:

Met Thr Tyr Leu Met Asn Asn Tyr Ala Arg Leu Pro Val Lys Phe Val
 5 10 15
 Arg Gly Lys Gly Val Tyr Leu Tyr Asp Glu Glu Gly Lys Glu Tyr Leu
 20 25 30
 Asp Phe Val Ser Gly Ile Gly Val Asn Ser Leu Gly His Ala Tyr Pro
 35 40 45
 Lys Leu Thr Glu Ala Leu Lys Glu Gln Val Glu Lys Leu Leu His Val
 50 55 60
 Ser Asn Leu Tyr Glu Asn Pro Trp Gln Glu Glu Leu Ala His Lys Leu
 65 70 75 80

(xi) SEQUENCE DESCRIPTION: SEQ ID NO:29:

Met Arg Lys Leu Ala Glu Arg Ala Gln Lys Leu Ser Pro Ser Pro Thr
5 10 15

Leu Ser Val Asp Thr Lys Ala Lys Glu Leu Leu Arg Gln Gly Glu Arg
20 25 30

Val Ile Asn Phe Gly Ala Gly Glu Pro Asp Phe Asp Thr Pro Glu His
35 40 45

Ile Lys Glu Ala Ala Lys Arg Ala Leu Asp Gln Gly Phe Thr Lys Tyr
50 55 60

Thr Pro Val Ala Gly Ile Leu Pro Leu Arg Glu Ala Ile Cys Glu Lys
65 70 75 80

Leu Tyr Arg Asp Asn Gln Leu Glu Tyr Ser Pro Asn Glu Ile Val Val
85 90 95

Ser Cys Gly Ala Lys His Ser Ile Phe Asn Ala Leu Gln Val Leu Leu
100 105 110

Asp Pro Gly Asp Glu Val Ile Ile Pro Val Pro Tyr Trp Thr Ser Tyr
115 120 125

Pro Glu Gln Val Lys Leu Ala Gly Gly Val Pro Val Phe Val Pro Thr
130 135 140

Ser Pro Glu Asn Asp Phe Lys Leu Arg Pro Glu Asp Leu Arg Ala Ala
145 150 155 160

Val Thr Pro Arg Thr Arg Leu Leu Ile Leu Asn Ser Pro Ala Asn Pro
165 170 175

Thr Gly Thr Val Tyr Arg Arg Glu Glu Leu Ile Gly Leu Ala Glu Val
180 185 190

Ala Leu Glu Ala Asp Leu Trp Ile Leu Ser Asp Glu Ile Tyr Glu Lys
195 200 205

Leu Ile Tyr Asp Gly Met Glu His Val Ser Ile Ala Ala Leu Asp Pro
210 215 220

Glu Val Lys Lys Arg Thr Ile Val Val Asn Gly Val Ser Lys Ala Tyr
225 230 235 240

Ala Met Thr Gly Trp Arg Ile Gly Tyr Ala Ala Ala Pro Arg Pro Ile
245 250 255

Ala Gln Ala Met Thr Asn Leu Gln Ser His Ser Thr Ser Asn Pro Thr
260 265 270

Ser Val Ala Gln Ala Ala Ala Leu Ala Ala Leu Lys Gly Pro Gln Glu
275 280 285

Pro Val Glu Asn Met Arg Arg Ala Phe Gln Lys Arg Arg Asp Phe Ile
290 295 300

Trp Gln Tyr Leu Asn Ser Leu Pro Gly Val Arg Cys Pro Lys Pro Leu
305 310 315 320

Gly Ala Phe Tyr Val Phe Pro Glu Val Glu Arg Ala Phe Gly Pro Pro
325 330 335

Ser Lys Arg Thr Gly Asn Thr Thr Ala Ser Asp Leu Ala Leu Phe Leu
340 345 350

Leu Glu Glu Ile Lys Val Ala Thr Val Ala Gly Ala Ala Phe Gly Asp
355 360 365

Asp Arg Tyr Leu Arg Phe Ser Tyr Ala Leu Arg Leu Glu Asp Ile Glu
370 375 380

Glu Gly Met Gln Arg Phe Lys Glu Leu Ile Glu Ala Ala Leu
385 390 395

(2) INFORMATION FOR SEQ ID NO:30:

(i) SEQUENCE CHARACTERISTICS

(A) LENGTH: 592 AMINO ACIDS

(B) TYPE: AMINO ACID

(D) TOPOLOGY: LINEAR

(ii) MOLECULE TYPE: PROTEIN

(xi) SEQUENCE DESCRIPTION: SEQ ID NO:30:

Met Cys Gly Ile Val Gly Tyr Val Gly Arg Asp Leu Ala Leu Pro Ile
5 10 15

Val Leu Gly Ala Leu Glu Arg Leu Glu Tyr Arg Gly Tyr Asp Ser Ala
20 25 30

Gly Val Ala Leu Ile Glu Asp Gly Lys Leu Ile Val Glu Lys Lys Lys
35 40 45

Gly Lys Ile Arg Glu Leu Val Lys Ala Leu Trp Gly Lys Asp Tyr Lys
50 55 60

Ala Lys Thr Gly Ile Gly His Thr Arg Trp Ala Thr His Gly Lys Pro
65 70 75 80

Thr Asp Glu Asn Ala His Pro His Thr Asp Glu Lys Gly Glu Phe Ala
85 90 95

Val Val His Asn Gly Ile Ile Glu Asn Tyr Leu Glu Leu Lys Glu Glu
100 105 110

Leu Lys Lys Glu Gly Val Lys Phe Arg Ser Glu Thr Asp Thr Glu Val
115 120 125

Ile Ala His Leu Ile Ala Lys Asn Tyr Arg Gly Asp Leu Leu Glu Ala
130 135 140

Val Leu Lys Thr Val Lys Lys Leu Lys Gly Ala Phe Ala Phe Ala Val
145 150 155 160

Ile Thr Val His Glu Pro Asn Arg Leu Ile Gly Val Lys Gln Gly Ser
165 170 175

Pro Leu Ile Val Gly Leu Gly Glu Gly Glu Asn Phe Leu Ala Ser Asp
180 185 190

Ile Pro Ala Ile Leu Pro Tyr Thr Lys Lys Ile Ile Val Leu Asp Asp
 195 200 205
 Gly Glu Ile Ala Asp Leu Thr Pro Asp Thr Val Asn Ile Tyr Asn Phe
 210 215 220
 Glu Gly Glu Pro Val Ser Lys Glu Val Met Ile Thr Pro Trp Asp Leu
 225 230 235 240
 Val Ser Ala Glu Lys Gly Gly Phe Lys His Phe Met Leu Lys Glu Ile
 245 250 255
 Tyr Glu Gln Pro Lys Ala Ile Asn Asp Thr Leu Lys Gly Phe Leu Ser
 260 265 270
 Thr Glu Asp Ala Ile Pro Phe Lys Leu Lys Asp Phe Arg Arg Val Leu
 275 280 285
 Ile Ile Ala Cys Gly Thr Ser Tyr His Ala Gly Phe Val Gly Lys Tyr
 290 295 300
 Trp Ile Glu Arg Phe Ala Gly Val Pro Thr Glu Val Ile Tyr Ala Ser
 305 310 315 320
 Glu Phe Arg Tyr Ala Asp Val Pro Val Ser Asp Lys Asp Ile Val Ile
 325 330 335
 Gly Ile Ser Gln Ser Gly Glu Thr Ala Asp Thr Lys Phe Ala Leu Gln
 340 345 350
 Ser Ala Lys Glu Lys Gly Ala Phe Thr Val Gly Leu Val Asn Val Val
 355 360 365
 Gly Ser Ala Ile Asp Arg Glu Ser Asp Phe Ser Leu His Thr His Ala
 370 375 380
 Gly Pro Glu Ile Gly Val Ala Ala Thr Lys Thr Phe Thr Ala Gln Phe
 385 390 395 400
 Thr Ala Leu Tyr Ala Leu Ser Val Arg Glu Ser Glu Glu Arg Glu Asn
 405 410 415
 Leu Ile Arg Leu Leu Glu Lys Val Pro Ser Leu Val Glu Gln Thr Leu
 420 425 430
 Asn Thr Ala Glu Glu Val Glu Lys Val Ala Glu Lys Tyr Met Lys Lys
 435 440 445
 Lys Asn Met Leu Tyr Leu Gly Arg Tyr Leu Asn Tyr Pro Ile Ala Leu
 450 455 460
 Glu Gly Ala Leu Lys Leu Lys Glu Ile Ser Tyr Ile His Ala Glu Gly
 465 470 475 480
 Tyr Pro Ala Gly Glu Met Lys His Gly Pro Ile Ala Leu Ile Asp Glu
 485 490 495
 Asn Met Pro Val Val Val Ile Ala Pro Lys Asp Arg Val Tyr Glu Lys
 500 505 510
 Ile Leu Ser Asn Val Glu Glu Val Leu Ala Arg Lys Gly Arg Val Ile
 515 520 525

Ser Val Gly Phe Lys Gly Asp Glu Thr Leu Lys Ser Lys Ser Glu Ser
530 535 540

Val Met Glu Ile Pro Lys Ala Glu Glu Pro Ile Thr Pro Phe Leu Thr
545 550 555 560

Val Ile Pro Leu Gln Leu Phe Ala Tyr Phe Ile Ala Ser Lys Leu Gly
565 570 575

Leu Asp Val Asp Gln Pro Arg Asn Leu Ala Lys Thr Val Thr Val Glu
580 585 590

(2) INFORMATION FOR SEQ ID NO:31:

(i) SEQUENCE CHARACTERISTICS

(A) LENGTH: 354 AMINO ACIDS

(B) TYPE: AMINO ACID

(D) TOPOLOGY: LINEAR

(ii) MOLECULE TYPE: PROTEIN

(xi) SEQUENCE DESCRIPTION: SEQ ID NO:31:

Met Ile Pro Gln Arg Ile Lys Glu Leu Glu Ala Tyr Lys Thr Glu Val
5 10 15

Thr Pro Ala Ser Val Arg Leu Ser Ser Asn Glu Phe Pro Tyr Asp Phe
20 25 30

Pro Glu Glu Ile Lys Gln Arg Ala Leu Glu Glu Leu Lys Lys Val Pro
35 40 45

Leu Asn Lys Tyr Pro Asp Pro Glu Ala Lys Glu Leu Lys Ala Val Leu
50 55 60

Ala Asp Phe Phe Gly Val Lys Glu Glu Asn Leu Val Leu Gly Asn Gly
65 70 75 80

Ser Asp Glu Leu Ile Tyr Tyr Leu Ser Ile Ala Ile Gly Glu Leu Tyr
85 90 95

Ile Pro Val Tyr Ile Pro Val Pro Thr Phe Pro Met Tyr Glu Ile Ser
100 105 110

Ala Lys Val Leu Gly Arg Pro Leu Val Lys Val Gln Leu Asp Glu Asn
115 120 125

Phe Asp Ile Asp Leu Glu Arg Ser Ile Glu Leu Ile Glu Lys Glu Lys
130 135 140

Pro Val Leu Gly Tyr Phe Ala Tyr Pro Asn Asn Pro Thr Gly Asn Leu
145 150 155 160

Phe Ser Arg Gly Lys Ile Glu Glu Ile Arg Asn Arg Gly Val Phe Cys
165 170 175

Val Ile Asp Glu Ala Tyr Tyr His Tyr Ser Gly Glu Thr Phe Leu Glu
180 185 190

Asp Ala Leu Lys Arg Glu Asp Thr Val Val Leu Arg Thr Leu Ser Lys
195 200 205

Ile Gly Met Ala Ser Leu Arg Val Gly Ile Leu Ile Gly Lys Gly Glu
210 215 220

Ile Val Ser Glu Ile Asn Lys Val Arg Leu Pro Phe Asn Val Thr Tyr
225 230 235 240

Pro Ser Gln Val Met Ala Lys Val Leu Leu Thr Glu Gly Arg Glu Phe
245 250 255

Leu Met Glu Lys Ile Gln Glu Val Val Thr Glu Arg Glu Arg Met Tyr
260 265 270

Asp Glu Met Lys Lys Ile Glu Gly Val Glu Val Phe Pro Ser Lys Ala
275 280 285

Asn Phe Leu Leu Phe Arg Thr Pro Tyr Pro Ala His Glu Val Tyr Gln
290 295 300

Glu Leu Leu Lys Arg Asp Val Leu Val Arg Asn Val Ser Tyr Met Glu
305 310 315 320

Gly Leu Gln Lys Cys Leu Arg Val Ser Val Gly Lys Pro Glu Glu Asn
325 330 335

Asn Lys Phe Leu Glu Ala Leu Glu Glu Ser Ile Lys Ser Leu Ser Ser
340 345 350

Ser Leu

- (2) INFORMATION FOR SEQ ID NO:32:
- (i) SEQUENCE CHARACTERISTICS
(A) LENGTH: 303 AMINO ACIDS
(B) TYPE: AMINO ACID
(D) TOPOLOGY: LINEAR
- (ii) MOLECULE TYPE: PROTEIN
- (xi) SEQUENCE DESCRIPTION: SEQ ID NO:32:

Met Lys Pro Tyr Ala Lys Tyr Ile Trp Leu Asp Gly Arg Ile Leu Lys
5 10 15

Trp Glu Asp Ala Lys Ile His Val Leu Thr His Ala Leu His Tyr Gly
20 25 30

Thr Ser Ile Phe Glu Gly Ile Arg Gly Tyr Trp Asn Gly Asp Asn Leu
35 40 45

Leu Val Phe Arg Leu Glu Glu His Ile Asp Arg Met Tyr Arg Ser Ala
50 55 60

Lys Ile Leu Gly Ile Asn Ile Pro Tyr Thr Arg Glu Glu Val Arg Gln
65 70 75 80

Ala Val Leu Glu Thr Ile Lys Ala Asn Asn Phe Arg Glu Asp Val Tyr
85 90 95

Ile Arg Pro Val Ala Phe Val Ala Ser Gln Thr Val Thr Leu Asp Ile
100 105 110

Arg Asn Leu Glu Val Ser Leu Ala Val Ile Val Phe Pro Phe Gly Lys
 115 120 125
 Tyr Leu Ser Pro Asn Gly Ile Lys Ala Thr Ile Val Ser Trp Arg Arg
 130 135 140
 Val His Asn Thr Met Leu Pro Val Met Ala Lys Ile Gly Gly Ile Tyr
 145 150 155 160
 Val Asn Ser Val Leu Ala Leu Val Glu Ala Arg Ser Arg Gly Phe Asp
 165 170 175
 Glu Ala Leu Leu Met Asp Val Asn Gly Tyr Val Val Glu Gly Ser Gly
 180 185 190
 Glu Asn Ile Phe Ile Val Arg Gly Gly Arg Leu Phe Thr Pro Pro Val
 195 200 205
 His Glu Ser Ile Leu Glu Gly Ile Thr Arg Asp Thr Val Ile Lys Leu
 210 215 220
 Ser Gly Asp Val Gly Leu Arg Val Glu Glu Lys Pro Ile Thr Arg Glu
 225 230 235 240
 Glu Val Tyr Thr Ala Asp Glu Val Phe Leu Val Gly Thr Ala Ala Glu
 245 250 255
 Ile Thr Pro Val Val Glu Val Asp Gly Arg Thr Ile Gly Thr Gly Lys
 260 265 270
 Pro Gly Pro Ile Thr Thr Lys Ile Ala Glu Leu Tyr Ser Asn Val Val
 275 280 285
 Arg Gly Lys Val Glu Lys Tyr Leu Asn Trp Ile Thr Pro Val Tyr
 290 295 300

(2) INFORMATION FOR SEQ ID NO:33:

- (i) SEQUENCE CHARACTERISTICS
 - (A) LENGTH: 52 NUCLEOTIDES
 - (B) TYPE: NUCLEIC ACID
 - (C) STRANDEDNESS: SINGLE
 - (D) TOPOLOGY: LINEAR

(ii) MOLECULE TYPE: cDNA

(xi) SEQUENCE DESCRIPTION: SEQ ID NO:33:

CCGAGAATTC ATTAAAGAGG AGAAATTAAC TATGGCAGTC AAAGTGCGGC CT

52

(2) INFORMATION FOR SEQ ID NO:34:

- (i) SEQUENCE CHARACTERISTICS
 - (A) LENGTH: 31 NUCLEOTIDES
 - (B) TYPE: NUCLEIC ACID
 - (C) STRANDEDNESS: SINGLE
 - (D) TOPOLOGY: LINEAR

- (ii) MOLECULE TYPE: cDNA
- (xi) SEQUENCE DESCRIPTION: SEQ ID NO:34:

CGGAGGATCC TTATCCAAAG CTTCCAGGAA G

31

(2) INFORMATION FOR SEQ ID NO:35:

- (i) SEQUENCE CHARACTERISTICS
- (A) LENGTH: 1,092 NUCLEOTIDES
 - (B) TYPE: NUCLEIC ACID
 - (C) STRANDEDNESS: SINGLE
 - (D) TOPOLOGY: LINEAR

- (ii) MOLECULE TYPE: GENOMIC DNA

- (xi) SEQUENCE DESCRIPTION: SEQ ID NO:35:

ATG GCA GTC AAA GTG CGG CCT GAG CTC AGC CAG GTG GAG ATC TAC CGT	48
Met Ala Val Lys Val Arg Pro Glu Leu Ser Gln Val Glu Ile Tyr Arg	
5 10 15	
CCC GGC AAA CCC ATC GAA GAG GTA AAG AAG GAG CTG GGG CTG GAG GAG	96
Pro Gly Lys Pro Ile Glu Glu Val Lys Lys Glu Leu Gly Leu Glu Glu	
20 25 30	
GTA GTC AAG CTG GCC TCC AAC GAG AAC CCT CTG GGA CCT TCT CCC AAG	144
Val Val Lys Leu Ala Ser Asn Glu Asn Pro Leu Gly Pro Ser Pro Lys	
35 40 45	
GCC GTG GCG GCG CTG GAG GGA CTG GAC CAC TGG CAC CTT TAC CCA GAA	192
Ala Val Ala Ala Leu Glu Gly Leu Asp His Trp His Leu Tyr Pro Glu	
50 55 60	
GGC TCA AGC TAT GAG CTA CGG CAG GCG CTG GGT AAG AAA CTG GAG ATA	240
Gly Ser Ser Tyr Glu Leu Arg Gln Ala Leu Gly Lys Lys Leu Glu Ile	
65 70 75 80	
GAC CCG GAC AGC ATC ATC GTG GGT TGC GGC TCA AGC GAA GTC ATC CAG	288
Asp Pro Asp Ser Ile Ile Val Gly Cys Gly Ser Ser Glu Val Ile Gln	
85 90 95	
ATG CTC TCT TTG GCC CTG CTG GCG CCC GGC GAC GAG GTG GTC ATC CCT	336
Met Leu Ser Leu Ala Leu Leu Ala Pro Gly Asp Glu Val Val Ile Pro	
100 105 110	
GTG CCT ACC TTT CCC CGC TAT GAG CCC CTG GCA CGG CTC ATG GGG GCT	384
Val Pro Thr Phe Pro Arg Tyr Glu Pro Leu Ala Arg Leu Met Gly Ala	
115 120 125	
AAT CCC GTA AAA GTT CCC TTG AAG GAC TAC CGC ATC GAT GTG GAG GCA	432
Asn Pro Val Lys Val Pro Leu Lys Asp Tyr Arg Ile Asp Val Glu Ala	
130 135 140	
GTG GCC CGA GCC CTT TCC CCC CGT ACC AAG CTG GTC TAC CTA TGC AAC	480
Val Ala Arg Ala Leu Ser Pro Arg Thr Lys Leu Val Tyr Leu Cys Asn	
145 150 155 160	
CCC AAC AAC CCC ACC GGG ACC ATC GTC ACC CGG GAG GAG GTG GAG TGG	528
Pro Asn Asn Pro Thr Gly Thr Ile Val Thr Arg Glu Glu Val Glu Trp	
165 170 175	

TTC	TTG	GAA	AAG	GCG	GGG	GAG	GGG	GTT	CTC	ACC	GTG	CTG	GAC	GAG	GCC	576
Phe	Leu	Glu	Lys	Ala	Gly	Glu	Gly	Val	Leu	Thr	Val	Leu	Asp	Glu	Ala	
			180					185					190			
TAC	TGC	GAG	TAC	GTG	ACC	AGC	CCC	GCC	TAC	CCT	GAT	GGG	CTC	GAT	TTC	624
Tyr	Cys	Glu	Tyr	Val	Thr	Ser	Pro	Ala	Tyr	Pro	Asp	Gly	Leu	Asp	Phe	
		195					200					205				
CTG	CGC	CGG	GGC	TAC	AAT	GTG	GTG	GTG	CTG	CGC	ACC	TTC	TCC	AAG	ATC	672
Leu	Arg	Arg	Gly	Tyr	Asn	Val	Val	Val	Leu	Arg	Thr	Phe	Ser	Lys	Ile	
	210					215					220					
TAC	GGG	CTG	GCC	GGG	CTG	CGC	ATA	GGG	TAC	GGT	GTG	GCG	GAC	AGG	GAG	720
Tyr	Gly	Leu	Ala	Gly	Leu	Arg	Ile	Gly	Tyr	Gly	Val	Ala	Asp	Arg	Glu	
225					230					235					240	
CTG	GTG	GCG	GAA	CTG	CAC	CGG	GTG	CGG	GAG	CCT	TTC	AAT	GTC	AGT	TCC	768
Leu	Val	Ala	Glu	Leu	His	Arg	Val	Arg	Glu	Pro	Phe	Asn	Val	Ser	Ser	
			245						250					255		
GCT	GCT	CAG	ATA	GCC	GCC	CTG	GCC	GCC	CTG	GAA	GAC	GAA	GAG	TTC	GTG	816
Ala	Ala	Gln	Ile	Ala	Ala	Leu	Ala	Ala	Leu	Glu	Asp	Glu	Glu	Phe	Val	
			260					265					270			
GCG	CTT	TCG	CGC	CAG	GTC	AAC	GAA	GAA	GGG	AAG	GTT	TTT	CTC	TAC	CGA	864
Ala	Leu	Ser	Arg	Gln	Val	Asn	Glu	Glu	Gly	Lys	Val	Phe	Leu	Tyr	Arg	
		275					280						285			
GAA	CTG	GAG	AGG	CGG	GGG	ATC	GCC	TAC	GTG	CCC	ACC	GAG	GCC	AAC	TTC	912
Glu	Leu	Glu	Arg	Arg	Gly	Ile	Ala	Tyr	Val	Pro	Thr	Glu	Ala	Asn	Phe	
	290					295					300					
CTA	CTC	TTC	GAT	GCC	GGT	CGG	GAC	GAG	CAG	GAA	GTA	TTT	CGC	CGG	ATG	960
Leu	Leu	Phe	Asp	Ala	Gly	Arg	Asp	Glu	Gln	Glu	Val	Phe	Arg	Arg	Met	
305					310					315					320	
CTG	CGC	CAG	GGA	GTG	ATC	ATC	CGG	GNC	GGG	GTG	GGT	TAT	CCC	ACC	CAC	1008
Leu	Arg	Gln	Gly	Val	Ile	Ile	Arg	Xxx	Gly	Val	Gly	Tyr	Pro	Thr	His	
			325					330						335		
TTA	AGG	GTG	ACC	ATC	GGC	ACC	TTG	GAA	CAG	AAC	CAG	CGC	TTC	CTG	GAA	1056
Leu	Arg	Val	Thr	Ile	Gly	Thr	Leu	Glu	Gln	Asn	Gln	Arg	Phe	Leu	Glu	
			340				345						350			
GCT	TTG	GAT	AAG	GCT	CTA	GAG	CTT	AGG	GGG	GTT	TAA					1092
Ala	Leu	Asp	Lys	Ala	Leu	Glu	Leu	Arg	Gly	Val						
		355					360			363						

(2) INFORMATION FOR SEQ ID NO:36:

- (i) SEQUENCE CHARACTERISTICS
 (A) LENGTH: AMINO ACIDS
 (B) TYPE: NUCLEIC ACID
 (C) STRANDEDNESS: SINGLE
 (D) TOPOLOGY: LINEAR

(ii) MOLECULE TYPE: cDNA

(xi) SEQUENCE DESCRIPTION: SEQ ID NO:36:

Met Ala Val Lys Val Arg Pro Glu Leu Ser Gln Val Glu Ile Tyr Arg
 5 10 15
 Pro Gly Lys Pro Ile Glu Glu Val Lys Lys Glu Leu Gly Leu Glu Glu

[illegible]

(i) SEQUENCE CHARACTERISTICS
(A) LENGTH: 52 NUCLEOTIDES
(B) TYPE: NUCLEIC ACID
(C) STRANDEDNESS: SINGLE
(D) TOPOLOGY: LINEAR

(xi) SEQUENCE DESCRIPTION: SEQ ID NO:37:

52

(2) INFORMATION FOR SEQ ID NO:38:

- (i) SEQUENCE CHARACTERISTICS
(A) LENGTH: 31 NUCLEOTIDES
(B) TYPE: NUCLEIC ACID
(C) STRANDEDNESS: SINGLE
(D) TOPOLOGY: LINEAR
- (ii) MOLECULE TYPE: cDNA
- (xi) SEQUENCE DESCRIPTION: SEQ ID NO:38:

CGGAGGATCC TTAGATCTCT TCAAGGGCTT T

31

(2) INFORMATION FOR SEQ ID NO:39:

- (i) SEQUENCE CHARACTERISTICS
(A) LENGTH: 1,085 NUCLEOTIDES
(B) TYPE: NUCLEIC ACID
(C) STRANDEDNESS: SINGLE
(D) TOPOLOGY: LINEAR
- (ii) MOLECULE TYPE: nucleic acid
- (xi) SEQUENCE DESCRIPTION: SEQ ID NO:39:

ATG AGA AAA GGA CTT GCA AGT AGG GTA AGT CAC CTA AAA CCT TCC CCC	48
Met Arg Lys Gly Leu Ala Ser Arg Val Ser His Leu Lys Pro Ser Pro	
5 10 15	
ACG CTG ACC ATA ACC GCA AAA GCA AAA GAA TTA AGG GCT AAA GGA GTG	96
Thr Leu Thr Ile Thr Ala Lys Ala Lys Glu Leu Arg Ala Lys Gly Val	
20 25 30	
GAC GTT ATA GGT TTT GGA GCG GGA GAA CCT GAC TTC GAC ACA CCC GAC	144
Asp Val Ile Gly Phe Gly Ala Gly Glu Pro Asp Phe Asp Thr Pro Asp	
35 40 45	
TTC ATA AAG GAA GCC TGT ATA AGG GCT TTA AGG GAA GGA AAG ACC AAG	192
Phe Ile Lys Glu Ala Cys Ile Arg Ala Leu Arg Glu Gly Lys Thr Lys	
50 55 60	
TAC GCT CCC TCC GCG GGA ATA CCA GAG CTC AGA GAA GCT ATA GCT GAA	240
Tyr Ala Pro Ser Ala Gly Ile Pro Glu Leu Arg Glu Ala Ile Ala Glu	
65 70 75 80	
AAA CTA CTG AAA GAA AAC AAA GTT GAG TAC AAA CCT TCA GAG ATA GTC	288
Lys Leu Leu Lys Glu Asn Lys Val Glu Tyr Lys Pro Ser Glu Ile Val	
85 90 95	
GTT TCC GCA GGA GCG AAA ATG GTT CTC TTC CTC ATA TTC ATG GCT ATA	336
Val Ser Ala Gly Ala Lys Met Val Leu Phe Leu Ile Phe Met Ala Ile	
100 105 110	
CTG GAC GAA GGA GAC GAG GTT TTA CTA CCT AGC CCT TAC TGG GTA ACT	384
Leu Asp Glu Gly Asp Glu Val Leu Leu Pro Ser Pro Tyr Trp Val Thr	
115 120 125	
TAC CCC GAA CAG ATA AGG TTC TTC GGA GGG GTT CCC GTT GAG GTT CCT	432
Tyr Pro Glu Gln Ile Arg Phe Phe Gly Gly Val Pro Val Glu Val Pro	
130 135 140	
CTA AAG AAA GAG AAA GGA TTT CAA TTA AGT CTG GAA GAT GTG AAA GAA	480
Leu Lys Lys Glu Lys Gly Phe Gln Leu Ser Leu Glu Asp Val Lys Glu	
145 150 155 160	

GCG CTT TCG CGC CAG GTC AAC GAA GAA GGG AAG GTT TTT CTC TAC CGA	864
Ala Leu Ser Arg Gln Val Asn Glu Glu Gly Lys Val Phe Leu Tyr Arg	
275 280 285	
GAA CTG GAG AGG CGG GGG ATC GCC TAC GTG CCC ACC GAG GCC AAC TTC	912
Glu Leu Glu Arg Arg Gly Ile Ala Tyr Val Pro Thr Glu Ala Asn Phe	
290 295 300	
CTA CTC TTC GAT GCC GGT CGG GAC GAG CAG GAA GTA TTT CGC CGG ATG	960
Leu Leu Phe Asp Ala Gly Arg Asp Glu Gln Glu Val Phe Arg Arg Met	
305 310 315 320	
CTG CGC CAG GGA GTG ATC ATC CGG GNC GGG GTG GGT TAT CCC ACC CAC	1008
Leu Arg Gln Gly Val Ile Ile Arg Xxx Gly Val Gly Tyr Pro Thr His	
325 330 335	
TTA AGG GTG ACC ATC GGC ACC TTG GAA CAG AAC CAG CGC TTC CTG GAA	1056
Leu Arg Val Thr Ile Gly Thr Leu Glu Gln Asn Gln Arg Phe Leu Glu	
340 345 350	
GCT TTG GAT AAG GCT CTA GAG CTT AGG GGG GTT TAA	1092
Ala Leu Asp Lys Ala Leu Glu Leu Arg Gly Val	
355 360 363	

(2) INFORMATION FOR SEQ ID NO:36:

- (i) SEQUENCE CHARACTERISTICS
 (A) LENGTH: AMINO ACIDS
 (B) TYPE: NUCLEIC ACID
 (C) STRANDEDNESS: SINGLE
 (D) TOPOLOGY: LINEAR

(ii) MOLECULE TYPE: cDNA

(xi) SEQUENCE DESCRIPTION: SEQ ID NO:36:

Met	Ala	Val	Lys	Val	Arg	Pro	Glu	Leu	Ser	Gln	Val	Glu	Ile	Tyr	Arg
				5					10					15	
Pro	Gly	Lys	Pro	Ile	Glu	Glu	Val	Lys	Lys	Glu	Leu	Gly	Leu	Glu	Glu
			20					25					30		
Val	Val	Lys	Leu	Ala	Ser	Asn	Glu	Asn	Pro	Leu	Gly	Pro	Ser	Pro	Lys
		35					40					45			
Ala	Val	Ala	Ala	Leu	Glu	Gly	Leu	Asp	His	Trp	His	Leu	Tyr	Pro	Glu
		50				55				60					
Gly	Ser	Ser	Tyr	Glu	Leu	Arg	Gln	Ala	Leu	Gly	Lys	Lys	Leu	Glu	Ile
65				70						75				80	
Asp	Pro	Asp	Ser	Ile	Ile	Val	Gly	Cys	Gly	Ser	Ser	Glu	Val	Ile	Gln
				85				90						95	
Met	Leu	Ser	Leu	Ala	Leu	Leu	Ala	Pro	Gly	Asp	Glu	Val	Val	Ile	Pro
			100					105					110		

Val	Pro	Thr	Phe	Pro	Arg	Tyr	Glu	Pro	Leu	Ala	Arg	Leu	Met	Gly	Ala		
		115					120					125					
Asn	Pro	Val	Lys	Val	Pro	Leu	Lys	Asp	Tyr	Arg	Ile	Asp	Val	Glu	Ala		
	130					135					140						
Val	Ala	Arg	Ala	Leu	Ser	Pro	Arg	Thr	Lys	Leu	Val	Tyr	Leu	Cys	Asn		
145					150					155					160		
Pro	Asn	Asn	Pro	Thr	Gly	Thr	Ile	Val	Thr	Arg	Glu	Glu	Val	Glu	Trp		
				165					170					175			
Phe	Leu	Glu	Lys	Ala	Gly	Glu	Gly	Val	Leu	Thr	Val	Leu	Asp	Glu	Ala		
			180					185					190				
Tyr	Cys	Glu	Tyr	Val	Thr	Ser	Pro	Ala	Tyr	Pro	Asp	Gly	Leu	Asp	Phe		
	195						200					205					
Leu	Arg	Arg	Gly	Tyr	Asn	Val	Val	Val	Leu	Arg	Thr	Phe	Ser	Lys	Ile		
	210					215					220						
Tyr	Gly	Leu	Ala	Gly	Leu	Arg	Ile	Gly	Tyr	Gly	Val	Ala	Asp	Arg	Glu		
225					230					235					240		
Leu	Val	Ala	Glu	Leu	His	Arg	Val	Arg	Glu	Pro	Phe	Asn	Val	Ser	Ser		
				245					250					255			
Ala	Ala	Gln	Ile	Ala	Ala	Leu	Ala	Ala	Leu	Glu	Asp	Glu	Glu	Phe	Val		
			260					265					270				
Ala	Leu	Ser	Arg	Gln	Val	Asn	Glu	Glu	Gly	Lys	Val	Phe	Leu	Tyr	Arg		
		275					280					285					
Glu	Leu	Glu	Arg	Arg	Gly	Ile	Ala	Tyr	Val	Pro	Thr	Glu	Ala	Asn	Phe		
	290					295					300						
Leu	Leu	Phe	Asp	Ala	Gly	Arg	Asp	Glu	Gln	Glu	Val	Phe	Arg	Arg	Met		
305					310					315					320		
Leu	Arg	Gln	Gly	Val	Ile	Ile	Arg	Xxx	Gly	Val	Gly	Tyr	Pro	Thr	His		
				325					330					335			
Leu	Arg	Val	Thr	Ile	Gly	Thr	Leu	Glu	Gln	Asn	Gln	Arg	Phe	Leu	Glu		
			340					345					350				
Ala	Leu	Asp	Lys	Ala	Leu	Glu	Leu	Arg	Gly	Val							
		355					360			363							

(2) INFORMATION FOR SEQ ID NO:37:

(i) SEQUENCE CHARACTERISTICS

(A) LENGTH: 52 NUCLEOTIDES

- (B) TYPE: NUCLEIC ACID
- (C) STRANDEDNESS: SINGLE
- (D) TOPOLOGY: LINEAR

(ii) MOLECULE TYPE: cDNA

(xi) SEQUENCE DESCRIPTION: SEQ ID NO:37:

CCGAGAATTC ATTAAAGAGG AGAAATTAAC TATGAGAAAA GGACTTGCAA GT
52

(2) INFORMATION FOR SEQ ID NO:38:

- (i) SEQUENCE CHARACTERISTICS
 - (A) LENGTH: 31 NUCLEOTIDES
 - (B) TYPE: NUCLEIC ACID
 - (C) STRANDEDNESS: SINGLE
 - (D) TOPOLOGY: LINEAR

(ii) MOLECULE TYPE: cDNA

(xi) SEQUENCE DESCRIPTION: SEQ ID NO:38:

CGGAGGATCC TTAGATCTCT TCAAGGGCTT T
31

(2) INFORMATION FOR SEQ ID NO:39:

- (i) SEQUENCE CHARACTERISTICS
 - (A) LENGTH: 1,085 NUCLEOTIDES
 - (B) TYPE: NUCLEIC ACID
 - (C) STRANDEDNESS: SINGLE
 - (D) TOPOLOGY: LINEAR

(ii) MOLECULE TYPE: nucleic acid

(xi) SEQUENCE DESCRIPTION: SEQ ID NO:39:

ATG	AGA	AAA	GGA	CTT	GCA	AGT	AGG	GTA	AGT	CAC	CTA	AAA	CCT	TCC	CC	48
Met	Arg	Lys	Gly	Leu	Ala	Ser	Arg	Val	Ser	His	Leu	Lys	Pro	Ser	Pro	
				5					10						15	

ACG	CTG	ACC	ATA	ACC	GCA	AAA	GCA	AAA	GAA	TTA	AGG	GCT	AAA	GGA	GT	6
Thr	Leu	Thr	Ile	Thr	Ala	Lys	Ala	Lys	Glu	Leu	Arg	Ala	Lys	Gly	Val	
			20					25					30			

GAC	GTT	ATA	GGT	TTT	GGA	GCG	GGA	GAA	CCT	GAC	TTC	GAC	ACA	CCC	GA	4
Asp	Val	Ile	Gly	Phe	Gly	Ala	Gly	Glu	Pro	Asp	Phe	Asp	Thr	Pro	Asp	
			35				40					45				

TTC	ATA	AAG	GAA	GCC	TGT	ATA	AGG	GCT	TTA	AGG	GAA	GGA	AAG	ACC	AA	92
Phe	Ile	Lys	Glu	Ala	Cys	Ile	Arg	Ala	Leu	Arg	Glu	Gly	Lys	Thr	Lys	
	50					55					60					
TAC	GCT	CCC	TCC	GCG	GGA	ATA	CCA	GAG	CTC	AGA	GAA	GCT	ATA	GCT	GA	10
Tyr	Ala	Pro	Ser	Ala	Gly	Ile	Pro	Glu	Leu	Arg	Glu	Ala	Ile	Ala	Glu	
65					70					75					80	
AAA	CTA	CTG	AAA	GAA	AAC	AAA	GTT	GAG	TAC	AAA	CCT	TCA	GAG	ATA	GT	88
Lys	Leu	Leu	Lys	Glu	Asn	Lys	Val	Glu	Tyr	Lys	Pro	Ser	Glu	Ile	Val	
				85					90					95		
GTT	TCC	GCA	GGA	GCG	AAA	ATG	GTT	CTC	TTC	CTC	ATA	TTC	ATG	GCT	AB	86
Val	Ser	Ala	Gly	Ala	Lys	Met	Val	Leu	Phe	Leu	Ile	Phe	Met	Ala	Ile	
			100					105					110			
CTG	GAC	GAA	GGA	GAC	GAG	GTT	TTA	CTA	CCT	AGC	CCT	TAC	TGG	GTA	AG	84
Leu	Asp	Glu	Gly	Asp	Glu	Val	Leu	Leu	Pro	Ser	Pro	Tyr	Trp	Val	Thr	
		115					120					125				
TAC	CCC	GAA	CAG	ATA	AGG	TTC	TTC	GGA	GGG	GTT	CCC	GTT	GAG	GTT	CA	82
Tyr	Pro	Glu	Gln	Ile	Arg	Phe	Phe	Gly	Gly	Val	Pro	Val	Glu	Val	Pro	
	130					135					140					
CTA	AAG	AAA	GAG	AAA	GGA	TTT	CAA	TTA	AGT	CTG	GAA	GAT	GTG	AAA	GA	80
Leu	Lys	Lys	Glu	Lys	Gly	Phe	Gln	Leu	Ser	Leu	Glu	Asp	Val	Lys	Glu	
145					150					155					160	
AAG	GTT	ACG	GAG	AGA	ACA	AAA	GCT	ATA	GTC	ATA	AAC	TCT	CCG	AAC	AA	88
Lys	Val	Thr	Glu	Arg	Thr	Lys	Ala	Ile	Val	Ile	Asn	Ser	Pro	Asn	Asn	
				165					170					175		
CCC	ACT	GGT	GCT	GTT	TAC	GAA	GAG	GAG	GAA	CTT	AAG	AAA	ATA	GCG	GA	86
Pro	Thr	Gly	Ala	Val	Tyr	Glu	Glu	Glu	Glu	Leu	Lys	Lys	Ile	Ala	Glu	
			180				185						190			
TTT	TGC	GTG	GAG	AGG	GGC	ATT	TTC	ATA	ATT	TCC	GAT	GAG	TGC	TAT	GA	84
Phe	Cys	Val	Glu	Arg	Gly	Ile	Phe	Ile	Ile	Ser	Asp	Glu	Cys	Tyr	Glu	
	195					200						205				
TAC	TTC	GTT	TAC	GGT	GAT	GCA	AAA	TTT	GTT	AGC	CCT	GCC	TCT	TTC	TC	82
Tyr	Phe	Val	Tyr	Gly	Asp	Ala	Lys	Phe	Val	Ser	Pro	Ala	Ser	Phe	Ser	
	210					215					220					
GAT	GAA	GTA	AAG	AAC	ATA	ACC	TTC	ACG	GTA	AAC	GCC	TTT	TCG	AAG	AG	20
Asp	Glu	Val	Lys	Asn	Ile	Thr	Phe	Thr	Val	Asn	Ala	Phe	Ser	Lys	Ser	
225					230					235					240	
TAT	TCC	ATG	ACT	GGT	TGG	CGA	ATA	GGT	TAT	GTA	GCG	TGC	CCC	GAA	GA	88
Tyr	Ser	Met	Thr	Gly	Trp	Arg	Ile	Gly	Tyr	Val	Ala	Cys	Pro	Glu	Glu	
				245				250						255		
TAC	GCA	AAA	GTG	ATA	GCG	AGT	CTT	AAC	AGC	CAG	AGT	GTT	TCC	AAC	GB	16

Tyr	Ala	Lys	Val	Ile	Ala	Ser	Leu	Asn	Ser	Gln	Ser	Val	Ser	Asn	Val	
			260					265					270			
ACT	ACC	TTT	GCC	CAG	TAT	GGA	GCT	CTT	GAG	GCC	TTG	AAA	AAT	CCA	AAA	364
Thr	Thr	Phe	Ala	Gln	Tyr	Gly	Ala	Leu	Glu	Ala	Leu	Lys	Asn	Pro	Lys	
		275					280					285				
TCT	AAA	GAT	TTT	GTA	AAC	GAA	ATG	AGA	AAT	GCT	TTT	GAA	AGG	AGA	AGC	312
Ser	Lys	Asp	Phe	Val	Asn	Glu	Met	Arg	Asn	Ala	Phe	Glu	Arg	Arg	Arg	
	290					295					300					
GAT	ACG	GCT	GTA	GAA	GAG	CTT	TCT	AAA	ATT	CCA	GGT	ATG	GAT	GTG	GTA	350
Asp	Thr	Ala	Val	Glu	Glu	Leu	Ser	Lys	Ile	Pro	Gly	Met	Asp	Val	Val	
305					310					315					320	
AAA	CCC	GAA	GGT	GCC	TTT	TAC	ATA	TTT	CCG	GAC	TTC	TCC	GCT	TAC	GCT	308
Lys	Pro	Glu	Gly	Ala	Phe	Tyr	Ile	Phe	Pro	Asp	Phe	Ser	Ala	Tyr	Ala	
				325					330					335		
GAG	AAA	CTG	GGT	GGT	GAT	GTG	AAA	CTC	TCG	GAG	TTC	CTT	CTG	GAA	AAC	36
Glu	Lys	Leu	Gly	Gly	Asp	Val	Lys	Leu	Ser	Glu	Phe	Leu	Leu	Glu	Lys	
			340					345					350			
GCT	AAG	GTT	GCG	GTG	GTT	CCC	GGT	TCG	GCC	TTC	GGA	GCT	CCC	GGA	TTD	4
Ala	Lys	Val	Ala	Val	Val	Pro	Gly	Ser	Ala	Phe	Gly	Ala	Pro	Gly	Phe	
		355					360					365				
TTG	AGG	CTT	TCT	TAC	GCC	CTT	TCC	GAG	GAA	AGA	CTC	GTT	GAG	GGT	ATA	52
Leu	Arg	Leu	Ser	Tyr	Ala	Leu	Ser	Glu	Glu	Arg	Leu	Val	Glu	Gly	Ile	
	370					375					380					
AGG	AGA	ATA	AAG	AAA	GCC	CTT	GAA	GAG	ATC	TAA						1185
Arg	Arg	Ile	Lys	Lys	Ala	Leu	Glu	Glu	Ile							
385					390				394							

(2) INFORMATION FOR SEQ ID NO:40:

(i) SEQUENCE CHARACTERISTICS

- (A) LENGTH: 394 AMINO ACIDS
- (B) TYPE: AMINO ACID
- (D) TOPOLOGY: LINEAR

(ii) MOLECULE TYPE: polypeptide

(xi) SEQUENCE DESCRIPTION: SEQ ID NO:40:

Met	Arg	Lys	Gly	Leu	Ala	Ser	Arg	Val	Ser	His	Leu	Lys	Pro	Ser	Pro
				5					10					15	
Thr	Leu	Thr	Ile	Thr	Ala	Lys	Ala	Lys	Glu	Leu	Arg	Ala	Lys	Gly	Val

Asp	Thr	Ala	Val	Glu	Glu	Leu	Ser	Lys	Ile	Pro	Gly	Met	Asp	Val	Val
305					310					315					320
Lys	Pro	Glu	Gly	Ala	Phe	Tyr	Ile	Phe	Pro	Asp	Phe	Ser	Ala	Tyr	Ala
				325					330					335	
Glu	Lys	Leu	Gly	Gly	Asp	Val	Lys	Leu	Ser	Glu	Phe	Leu	Leu	Glu	Lys
			340					345					350		
Ala	Lys	Val	Ala	Val	Val	Pro	Gly	Ser	Ala	Phe	Gly	Ala	Pro	Gly	Phe
		355					360					365			
Leu	Arg	Leu	Ser	Tyr	Ala	Leu	Ser	Glu	Glu	Arg	Leu	Val	Glu	Gly	Ile
	370					375					380				
Arg	Arg	Ile	Lys	Lys	Ala	Leu	Glu	Glu	Ile						
385					390				394						

What Is Claimed Is:

1. An isolated polynucleotide comprising a member selected from the group consisting of:

- (a) a polynucleotide having at least a 70% identity to a polynucleotide encoding an enzyme comprising amino acid sequences set forth in SEQ ID NOS:25-32;
- (b) a polynucleotide which is complementary to the polynucleotide of (a); and
- (c) a polynucleotide comprising at least 15 consecutive bases of the polynucleotide of (a) or (b).

2. The polynucleotide of Claim 1 wherein the polynucleotide is DNA.

3. The polynucleotide of Claim 1 wherein the polynucleotide is RNA.

4. The polynucleotide of Claim 2 which encodes an enzyme comprising amino acids 1 to 414 of SEQ ID NO:25.

5. The polynucleotide of Claim 2 which encodes an enzyme comprising amino acids 1 to 373 of SEQ ID NO:26.

6. The polynucleotide of Claim 2 which encodes an enzyme comprising amino acids 1 to 453 of SEQ ID NO:27.

7. The polynucleotide of Claim 2 which encodes an enzyme comprising amino acids 1 to 343 of SEQ ID NO:28.

8. The polynucleotide of Claim 2 which encodes an enzyme comprising amino acids 1 to 398 of SEQ ID NO:29.

9. The polynucleotide of Claim 2 which encodes an enzyme comprising amino acids 1 to 592 of SEQ ID NO:30.

10. The polynucleotide of Claim 2 which encodes an enzyme comprising amino acids 1 to 354 of SEQ ID NO:31.

11. The polynucleotide of Claim 2 which encodes an enzyme comprising amino acids 1 to 303 of SEQ ID NO:32.

12. An isolated polynucleotide comprising a member selected from the group consisting of:

(a) a polynucleotide having at least a 70% identity to a polynucleotide encoding an enzyme expressed by the DNA contained in ATCC Deposit No. _____;

(b) a polynucleotide complementary to the polynucleotide of (a); and

(c) a polynucleotide comprising at least 15 consecutive bases of the polynucleotide of (a) and (b).

13. A vector comprising the DNA of Claim 2.

14. A host cell comprising the vector of Claim 13.

15. A process for producing a polypeptide comprising: expressing from the host cell of Claim 14 a polypeptide encoded by said DNA.

16. A process for producing a cell comprising: transforming or transfecting the cell with the vector of Claim 14 such that the cell expresses the polypeptide encoded by the DNA contained in the vector.

ABSTRACT

Thermostable transaminase and aminotransferase enzymes derived from various *ammonifex*, *aquifex* and ~~*pyrobaculum*~~ organisms are disclosed. The enzymes are produced from native or recombinant host cells and can be utilized in the pharmaceutical, agricultural and other industries.

[illegible]

FIGURE 3

ATG TGG GAA TTA GAC CCT AAA ACG CTC GAA AAG TGG GAC AAG GAG TAC	48
Met Trp Glu Leu Asp Pro Lys Thr Leu Glu Lys Trp Asp Lys Glu Tyr	
5 10 15	
TTC TGG CAT CCA TTT ACC CAG ATG AAA GTC TAC AGA GAA GAA GAA AAC	96
Phe Trp His Pro Phe Thr Gln Met Lys Val Tyr Arg Glu Glu Glu Asn	
20 25 30	
CTG ATA TTT GAA CGC GGA GAA GGC GTT TAC CTG TGG GAC ATA TAC GGC	144
Leu Ile Phe Glu Arg Gly Glu Gly Val Tyr Leu Trp Asp Ile Tyr Gly	
35 40 45	
AGG AAG TAT ATA GAT GCC ATA TCT TCC CTC TGG TGC AAC GTC CAC GGA	192
Arg Lys Tyr Ile Asp Ala Ile Ser Ser Leu Trp Cys Asn Val His Gly	
50 55 60	
CAT AAC CAC CCT AAA CTG AAC AAC GCA GTT ATG AAA CAG CTC TGT AAG	240
His Asn His Pro Lys Leu Asn Asn Ala Val Met Lys Gln Leu Cys Lys	
65 70 75 80	
GTA GCT CAC ACA ACT ACT CTG GGA AGT TCC AAC GTT CCC GCC ATA CTC	288
Val Ala His Thr Thr Thr Leu Gly Ser Ser Asn Val Pro Ala Ile Leu	
85 90 95	
CTT GCA AAG AAG CTT GTA GAA ATT TCT CCT GAA GGA TTA AAC AAG GTC	336
Leu Ala Lys Lys Leu Val Glu Ile Ser Pro Glu Gly Leu Asn Lys Val	
100 105 110	
TTT TAC TCC GAA GAC GGT GCG GAA GCA GTA GAG ATA GCG ATA AAG ATG	384
Phe Tyr Ser Glu Asp Gly Ala Glu Ala Val Glu Ile Ala Ile Lys Met	
115 120 125	
GCT TAT CAC TAC TGG AAG AAC AAG GGA GTT AAA GGG AAA AAC GTT TTC	432
Ala Tyr His Tyr Trp Lys Asn Lys Gly Val Lys Gly Lys Asn Val Phe	
130 135 140	
ATA ACG CTT TCC GAA GCC TAC CAC GGG GAT ACT GTA GGA GCG GTT AGC	480
Ile Thr Leu Ser Glu Ala Tyr His Gly Asp Thr Val Gly Ala Val Ser	
145 150 155 160	
GTA GGG GGT ATA GAA CTC TTC CAC GGA ACT TAT AAA GAT CTC CTT TTC	528
Val Gly Gly Ile Glu Leu Phe His Gly Thr Tyr Lys Asp Leu Leu Phe	
165 170 175	
AAG ACT ATA AAA CTC CCA TCT CCT TAC CTG TAC TGC AAG GAA AAG TAC	576
Lys Thr Ile Lys Leu Pro Ser Pro Tyr Leu Tyr Cys Lys Glu Lys Tyr	
180 185 190	
GGG GAA CTC TGC CCT GAG TGC ACG GCA GAT TTA TTA AAA CAA CTG GAA	624
Gly Glu Leu Cys Pro Glu Cys Thr Ala Asp Leu Leu Lys Gln Leu Glu	
195 200 205	
GAT ATC CTG AAG TCG CGG GAA GAT ATC GTT GCG GTC ATT ATG GAA GCG	672
Asp Ile Leu Lys Ser Arg Glu Asp Ile Val Ala Val Ile Met Glu Ala	
210 215 220	
GGA ATT CAG GCA GCC GCG GGA ATG CTC CCC TTC CCT CCG GGA TTT TTG	720
Gly Ile Gln Ala Ala Ala Gly Met Leu Pro Phe Pro Pro Gly Phe Leu	
225 230 235 240	

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AAA GGC GTA AGG GAG CTT ACG AAG AAA TAC GAC ACT TTA ATG ATA GTT Lys Gly Val Arg Glu Leu Thr Lys Lys Tyr Asp Thr Leu Met Ile Val 245 250 255	768
GAC GAG GTT GCC ACG GGA TTT GGC AGG ACG GGA ACG ATG TTT TAC TGT Asp Glu Val Ala Thr Gly Phe Gly Arg Thr Gly Thr Met Phe Tyr Cys 260 265 270	816
GAG CAG GAA GGA GTC AGT CCG GAC TTT ATG TGT CTA GGT AAG GGT ATA Glu Gln Glu Gly Val Ser Pro Asp Phe Met Cys Leu Gly Lys Gly Ile 275 280 285	864
ACC GGA GGG TAC CTC CCG CTT GCT GCG ACA CTC ACA ACG GAC GAG GTG Thr Gly Gly Tyr Leu Pro Leu Ala Ala Thr Leu Thr Thr Asp Glu Val 290 295 300	912
TTC AAT GCC TTT TTA GGT GAG TTC GGG GAG GCA AAG CAC TTT TAC CAC Phe Asn Ala Phe Leu Gly Glu Phe Gly Glu Ala Lys His Phe Tyr His 305 310 315 320	960
GGG CAC ACC TAC ACT GGA AAT AAC CTC GCC TGT TCC GTT GCA CTC GCA Gly His Thr Tyr Thr Gly Asn Asn Leu Ala Cys Ser Val Ala Leu Ala 325 330 335	1008
AAC TTA GAA GTT TTT GAG GAA GAA AGA ACT TTA GAG AAG CTC CAA CCA Asn Leu Glu Val Phe Glu Glu Glu Arg Thr Leu Glu Lys Leu Gln Pro 340 345 350	1056
AAG ATA AAG CTT TTA AAG GAA AGG CTT CAG GAG TTC TGG GAA CTC AAG Lys Ile Lys Leu Leu Lys Glu Arg Leu Gln Glu Phe Trp Glu Leu Lys 355 360 365	1104
CAC GTT GGA GAT GTT AGA CAG CTA GGT TTT ATG GCT GGA ATA GAG CTG His Val Gly Asp Val Arg Gln Leu Gly Phe Met Ala Gly Ile Glu Leu 370 375 380	1152
GTG AAG GAC AAA GAA AAG GGA GAA CCT TTC CCT TAC GGT GAA AGG ACG Val Lys Asp Lys Glu Lys Gly Glu Pro Phe Pro Tyr Gly Glu Arg Thr 385 390 395 400	1200
GGA TTT AAG GTG GCT TAC AAG TGC AGG GAA AAA GGG GTG TTT TTG AGA Gly Phe Lys Val Ala Tyr Lys Cys Arg Glu Lys Gly Val Phe Leu Arg 405 410 415	1245
CCG CTC GGA GAC GTT ATG GTA TTG ATG ATG CCT CTT GTA ATA GAG GAA Pro Leu Gly Asp Val Met Val Leu Met Met Pro Leu Val Ile Glu Glu 420 425 430	1293
GAC GAA ATG AAC TAC GTT ATT GAT ACA CTT AAA TGG GCA ATT AAA GAG Asp Glu Met Asn Tyr Val Ile Asp Thr Leu Lys Trp Ala Ile Lys Glu 435 440 445	1341
CTT GAA AAA GAG GTG TAG Leu Glu Lys Glu Val End 450	1359

FIGURE 4

ATG ACA TAC TTA ATG AAC AAT TAC GCA AGG TTG CCC GTA AAG TTT GTA	48
Met Thr Tyr Leu Met Asn Asn Tyr Ala Arg Leu Pro Val Lys Phe Val	
5 10 15	
AGG GGA AAA GGT GTT TAC CTG TAC GAT GAG GAA GGA AAG GAG TAT CTT	96
Arg Gly Lys Gly Val Tyr Leu Tyr Asp Glu Glu Gly Lys Glu Tyr Leu	
20 25 30	
GAC TTT GTC TCC GGT ATA GGC GTC AAC TCC CTC GGT CAC GCT TAC CCA	144
Asp Phe Val Ser Gly Ile Gly Val Asn Ser Leu Gly His Ala Tyr Pro	
35 40 45	
AAA CTC ACA GAA GCT CTA AAA GAA CAG GTT GAG AAA CTC CTC CAC GTT	192
Lys Leu Thr Glu Ala Leu Lys Glu Gln Val Glu Lys Leu Leu His Val	
50 55 60	
TCA AAT CTT TAC GAA AAC CCG TGG CAG GAA GAA CTG GCT CAC AAA CTT	240
Ser Asn Leu Tyr Glu Asn Pro Trp Gln Glu Glu Leu Ala His Lys Leu	
65 70 75 80	
GTA AAA CAC TTC TGG ACA GAA GGG AAG GTA TTT TTC GCA AAC AGC GGA	288
Val Lys His Phe Trp Thr Glu Gly Lys Val Phe Phe Ala Asn Ser Gly	
85 90 95	
ACG GAA AGT GTA GAG GCG GCT ATA AAG CTC GCA AGG AAG TAC TGG AGG	336
Thr Glu Ser Val Glu Ala Ala Ile Lys Leu Ala Arg Lys Tyr Trp Arg	
100 105 110	
GAT AAA GGA AAG AAC AAG TGG AAG TTT ATA TCC TTT GAA AAC TCT TTC	384
Asp Lys Gly Lys Asn Lys Trp Lys Phe Ile Ser Phe Glu Asn Ser Phe	
115 120 125	
CAC GGG AGA ACC TAC GGT AGC CTC TCC GCA ACG GGA CAG CCA AAG TTC	432
His Gly Arg Thr Tyr Gly Ser Leu Ser Ala Thr Gly Gln Pro Lys Phe	
130 135 140	
CAC AAA GGC TTT GAA CCT CTA GTT CCT GGA TTT TCT TAC GCA AAG CTG	480
His Lys Gly Phe Glu Pro Leu Val Pro Gly Phe Ser Tyr Ala Lys Leu	
145 150 155 160	
AAC GAT ATA GAC AGC GTT TAC AAA CTC CTA GAC GAG GAA ACC GCG GGG	528
Asn Asp Ile Asp Ser Val Tyr Lys Leu Leu Asp Glu Glu Thr Ala Gly	
165 170 175	
ATA ATT ATT GAA GTT ATA CAA GGA GAG GGC GGA GTA AAC GAG GCG AGT	576
Ile Ile Ile Glu Val Ile Gln Gly Glu Gly Gly Val Asn Glu Ala Ser	
180 185 190	
GAG GAT TTT CTA AGT AAA CTC CAG GAA ATT TGT AAA GAA AAA GAT GTG	624
Glu Asp Phe Leu Ser Lys Leu Gln Glu Ile Cys Lys Glu Lys Asp Val	
195 200 205	
CTC TTA ATT ATA GAC GAA GTG CAA ACG GGA ATA GGA AGG ACC GGG GAA	672
Leu Leu Ile Ile Asp Glu Val Gln Thr Gly Ile Gly Arg Thr Gly Glu	
210 215 220	
TTC TAC GCA TAT CAA CAC TTC AAT CTA AAA CCG GAC GTA ATT GCG CTT	720
Phe Tyr Ala Tyr Gln His Phe Asn Leu Lys Pro Asp Val Ile Ala Leu	
225 230 235 240	

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GCC ATG ACC GGT TGG CGC ATA GGT TAT GCT GCC GCT CCC CGG CCG ATA	768
Ala Met Thr Gly Trp Arg Ile Gly Tyr Ala Ala Ala Pro Arg Pro Ile	
245 250 255	
GCC CAG GCC ATG ACC AAC CTC CAA AGC CAC AGT ACC TCT AAC CCC ACT	816
Ala Gln Ala Met Thr Asn Leu Gln Ser His Ser Thr Ser Asn Pro Thr	
260 265 270	
TCC GTA GCC CAG GCG GCG GCG CTG GCC GCT CTG AAG GGG CCA CAA GAG	864
Ser Val Ala Gln Ala Ala Ala Leu Ala Ala Leu Lys Gly Pro Gln Glu	
275 280 285	
CCG GTG GAG AAC ATG CGC CGG GCT TTT CAA AAG CGG CGG GAT TTC ATC	912
Pro Val Glu Asn Met Arg Arg Ala Phe Gln Lys Arg Arg Asp Phe Ile	
290 295 300	
TGG CAG TAC CTA AAC TCC TTA CCC GGA GTG CGC TGC CCC AAA CCT TTA	960
Trp Gln Tyr Leu Asn Ser Leu Pro Gly Val Arg Cys Pro Lys Pro Leu	
305 310 315 320	
GGG GCC TTT TAC GTC TTT CCA GAA GTT GAG CGG GCT TTT GGG CCG CCG	1008
Gly Ala Phe Tyr Val Phe Pro Glu Val Glu Arg Ala Phe Gly Pro Pro	
325 330 335	
TCT AAA AGG ACG GGA AAT ACT ACC GCT AGC GAC CTG GCC CTT TTC CTC	1056
Ser Lys Arg Thr Gly Asn Thr Thr Ala Ser Asp Leu Ala Leu Phe Leu	
340 345 350	
CTG GAA GAG ATA AAA GTG GCC ACC GTG GCT GGG GCT GCC TTT GGG GAC	1104
Leu Glu Glu Ile Lys Val Ala Thr Val Ala Gly Ala Ala Phe Gly Asp	
355 360 365	
GAT CGC TAC CTG CGC TTT TCC TAC GCC CTG CGG CTG GAA GAT ATC GAA	1152
Asp Arg Tyr Leu Arg Phe Ser Tyr Ala Leu Arg Leu Glu Asp Ile Glu	
370 375 380	
GAG GGG ATG CAA CGG TTT AAA GAA TTG ATC GAA GCG GCA CTT TAA	1197
Glu Gly Met Gln Arg Phe Lys Glu Leu Ile Glu Ala Ala Leu End	
385 390 395	

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FIGURE 6

ATG TGC GGG ATA GTC GGA TAC GTA GGG AGG GAT TTA GCC CTT CCT ATA	48
Met Cys Gly Ile Val Gly Tyr Val Gly Arg Asp Leu Ala Leu Pro Ile	
5 10 15	
GTC CTC GGA GCT CTT GAG AGA CTC GAA TAC AGG GGT TAC GAC TCC GCG	96
Val Leu Gly Ala Leu Glu Arg Leu Glu Tyr Arg Gly Tyr Asp Ser Ala	
20 25 30	
GGA GTT GCC CTT ATA GAA GAC GGG AAA CTC ATA GTT GAA AAG AAG AAG	144
Gly Val Ala Leu Ile Glu Asp Gly Lys Leu Ile Val Glu Lys Lys Lys	
35 40 45	
GGA AAG ATA AGG GAA CTC GTT AAA GCG CTA TGG GGA AAG GAT TAC AAG	192
Gly Lys Ile Arg Glu Leu Val Lys Ala Leu Trp Gly Lys Asp Tyr Lys	
50 55 60	
GCT AAA ACG GGT ATA GGT CAC ACA CGC TGG GCA ACC CAC GGA AAG CCC	240
Ala Lys Thr Gly Ile Gly His Thr Arg Trp Ala Thr His Gly Lys Pro	
65 70 75 80	
ACG GAC GAG AAC GCC CAC CCC CAC ACC GAC GAA AAA GGT GAG TTT GCA	288
Thr Asp Glu Asn Ala His Pro His Thr Asp Glu Lys Gly Glu Phe Ala	
85 90 95	
GTA GTT CAC AAC GGG ATA ATA GAA AAC TAC TTA GAA CTA AAA GAG GAA	336
Val Val His Asn Gly Ile Ile Glu Asn Tyr Leu Glu Leu Lys Glu Glu	
100 105 110	
CTA AAG AAG GAA GGT GTA AAG TTC AGG TCC GAA ACA GAC ACA GAA GTT	384
Leu Lys Lys Glu Gly Val Lys Phe Arg Ser Glu Thr Asp Thr Glu Val	
115 120 125	
ATA GCC CAC CTC ATA GCG AAG AAC TAC AGG GGG GAC TTA CTG GAG GCC	432
Ile Ala His Leu Ile Ala Lys Asn Tyr Arg Gly Asp Leu Leu Glu Ala	
130 135 140	
GTT TTA AAA ACC GTA AAG AAA TTA AAG GGT GCT TTT GCC TTT GCG GTT	480
Val Leu Lys Thr Val Lys Lys Leu Lys Gly Ala Phe Ala Phe Ala Val	
145 150 155 160	
ATA ACG GTT CAC GAA CCA AAC AGA CTA ATA GGA GTG AAG CAG GGG AGT	528
Ile Thr Val His Glu Pro Asn Arg Leu Ile Gly Val Lys Gln Gly Ser	
165 170 175	
CCT TTA ATC GTC GGA CTC GGA GAA GGA GAA AAC TTC CTC GCT TCA GAT	576
Pro Leu Ile Val Gly Leu Gly Glu Gly Glu Asn Phe Leu Ala Ser Asp	
180 185 190	
ATT CCC GCA ATA CTT CCT TAC ACG AAA AAG ATT ATT GTT CTT GAT GAC	624
Ile Pro Ala Ile Leu Pro Tyr Thr Lys Lys Ile Ile Val Leu Asp Asp	
195 200 205	
GGG GAA ATA GCG GAC CTG ACT CCC GAC ACT GTG AAC ATT TAC AAC TTT	672
Gly Glu Ile Ala Asp Leu Thr Pro Asp Thr Val Asn Ile Tyr Asn Phe	
210 215 220	
GAG GGA GAG CCC GTT TCA AAG GAA GTA ATG ATT ACG CCC TGG GAT CTT	720
Glu Gly Glu Pro Val Ser Lys Glu Val Met Ile Thr Pro Trp Asp Leu	
225 230 235 240	

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[illegible]

FIGURE 7

ATG ATA CCC CAG AGG ATT AAG GAA CTT GAA GCT TAC AAG ACG GAG GTC Met Ile Pro Gln Arg Ile Lys Glu Leu Glu Ala Tyr Lys Thr Glu Val 5 10 15	48
ACT CCC GCC TCC GTC AGG CTT TCC TCT AAC GAA TTC CCC TAC GAC TTT Thr Pro Ala Ser Val Arg Leu Ser Ser Asn Glu Phe Pro Tyr Asp Phe 20 25 30	96
CCC GAG GAG ATA AAA CAA AGG GCC TTA GAA GAA TTA AAA AAG GTT CCC Pro Glu Glu Ile Lys Gln Arg Ala Leu Glu Glu Leu Lys Lys Val Pro 35 40 45	144
TTG AAC AAA TAC CCA GAC CCC GAA GCG AAA GAG TTA AAA GCG GTT CTT Leu Asn Lys Tyr Pro Asp Pro Glu Ala Lys Glu Leu Lys Ala Val Leu 50 55 60	192
GCG GAT TTT TTC GGC GTT AAG GAA GAA AAT TTA GTT CTC GGT AAC GGT Ala Asp Phe Phe Gly Val Lys Glu Glu Asn Leu Val Leu Gly Asn Gly 65 70 75 80	240
TCG GAC GAA CTC ATA TAC TAC CTC TCA ATA GCT ATA GGT GAA CTT TAC Ser Asp Glu Leu Ile Tyr Tyr Leu Ser Ile Ala Ile Gly Glu Leu Tyr 85 90 95	288
ATA CCC GTT TAC ATA CCT GTT CCC ACC TTT CCC ATG TAC GAG ATA AGT Ile Pro Val Tyr Ile Pro Val Pro Thr Phe Pro Met Tyr Glu Ile Ser 100 105 110	336
GCG AAA GTT CTC GGA AGA CCC CTC GTA AAG GTT CAA CTG GAC GAA AAC Ala Lys Val Leu Gly Arg Pro Leu Val Lys Val Gln Leu Asp Glu Asn 115 120 125	384
TTT GAT ATA GAC TTA GAA AGA AGT ATT GAA TTA ATA GAG AAA GAA AAA Phe Asp Ile Asp Leu Glu Arg Ser Ile Glu Leu Ile Glu Lys Glu Lys 130 135 140	432
CCC GTT CTC GGG TAC TTT GCT TAC CCA AAC AAC CCC ACG GGA AAC CTC Pro Val Leu Gly Tyr Phe Ala Tyr Pro Asn Asn Pro Thr Gly Asn Leu 145 150 155 160	480
TTT TCC AGG GGA AAG ATT GAG GAG ATA AGA AAC AGG GGT GTT TTC TGT Phe Ser Arg Gly Lys Ile Glu Glu Ile Arg Asn Arg Gly Val Phe Cys 165 170 175	528
GTA ATA GAC GAA GCC TAC TAT CAT TAC TCC GGA GAA ACC TTT CTG GAA Val Ile Asp Glu Ala Tyr Tyr His Tyr Ser Gly Glu Thr Phe Leu Glu 180 185 190	576
GAC GCG CTC AAA AGG GAA GAT ACG GTA GTT TTG AGG ACA CTT TCA AAA Asp Ala Leu Lys Arg Glu Asp Thr Val Val Leu Arg Thr Leu Ser Lys 195 200 205	624
ATC GGT ATG GCG AGT TTA AGG GTA GGG ATT TTA ATA GGG AAG GGG GAA Ile Gly Met Ala Ser Leu Arg Val Gly Ile Leu Ile Gly Lys Gly Glu 210 215 220	672
ATC GTC TCA GAA ATT AAC AAG GTG AGA CTC CCC TTC AAC GTG ACC TAC Ile Val Ser Glu Ile Asn Lys Val Arg Leu Pro Phe Asn Val Thr Tyr 225 230 235 240	720

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CCC TCT CAG GTG ATG GCA AAA GTT CTC CTC ACG GAG GGA AGA GAA TTC	768
Pro Ser Gln Val Met Ala Lys Val Leu Leu Thr Glu Gly Arg Glu Phe	
245 250 255	
CTA ATG GAA AAG ATA CAG GAG GTT GTA ACA GAG CGA GAA AGG ATG TAC	816
Leu Met Glu Lys Ile Gln Glu Val Val Thr Glu Arg Glu Arg Met Tyr	
260 265 270	
GAC GAA ATG AAG AAA ATA GAA GGA GTT GAG GTT TTT CCG AGT AAG GCT	864
Asp Glu Met Lys Lys Ile Glu Gly Val Glu Val Phe Pro Ser Lys Ala	
275 280 285	
AAC TTC TTG CTT TTC AGA ACG CCT TAC CCC GCC CAC GAG GTT TAT CAG	912
Asn Phe Leu Leu Phe Arg Thr Pro Tyr Pro Ala His Glu Val Tyr Gln	
290 295 300	
GAG CTA CTG AAA AGG GAT GTC CTC GTC AGG AAC GTA TCT TAC ATG GAA	960
Glu Leu Leu Lys Arg Asp Val Leu Val Arg Asn Val Ser Tyr Met Glu	
305 310 315 320	
GGA CTC CAA AAG TGC CTC AGG GTA AGC GTA GGG AAA CCG GAA GAA AAC	1008
Gly Leu Gln Lys Cys Leu Arg Val Ser Val Gly Lys Pro Glu Glu Asn	
325 330 335	
AAC AAG TTT CTG GAA GCA CTG GAG GAG AGT ATA AAA TCC CTT TCA AGC	1056
Asn Lys Phe Leu Glu Ala Leu Glu Glu Ser Ile Lys Ser Leu Ser Ser	
340 345 350	
TCT CTT TAA	1065
Ser Leu End	

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662050 733336

GAG Glu	GTG Val	TAT Tyr	ACA Thr	GCC Ala 245	GAC Asp	GAG Glu	GTG Val	TTT Phe 250	TTA Leu	GTA Val	GGA Gly	ACC Thr	GCC Ala 255	GCA Ala	GAG Glu	768
ATA Ile	ACG Thr	CCA Pro	GTG Val 260	GTG Val	GAG Glu	GTT Val	GAC Asp	GGC Gly 265	AGA Arg	ACA Thr	ATC Ile	GGC Gly 270	ACA Thr	GGC Gly	AAG Lys	816
CCG Pro	GGC Gly	CCC Pro 275	ATT Ile	ACG Thr	ACA Thr	AAA Lys	ATA Ile 280	GCT Ala	GAG Glu	CTG Leu	TAC Tyr	TCA Ser 285	AAC Asn	GTC Val	GTG Val	864
AGA	GGC	AAA	GTA	GAG	AAA	TAC	TTA	AAT	TGG	ATC	ACT	CCT	GTG	TAT	TAG	912
Arg 290	Gly	Lys	Val	Glu	Lys	Tyr 295	Leu	Asn	Trp	Ile	Thr 300	Pro	Val	Tyr	End	

FIGURE 9

Ammonifex degensii histidinol phosphate aminotransferase

1 ATG GCA GTC AAA GTG CGG CCT GAG CTC AGC CAG GTG GAG ATC TAC CGT CCC GGC AAA CCC 60 35
1 Met Ala Val Lys Val Arg Pro Glu Leu Ser Gln Val Glu Ile Tyr Arg Pro Gly Lys Pro 20 36

61 ATC GAA GAG GTA AAG AAG GAG CTG GGG CTG GAG GAG GTA GTC AAG CTG GCC TCC AAC GAG 120
21 Ile Glu Glu Val Lys Lys Glu Leu Gly Leu Glu Glu Val Val Lys Leu Ala Ser Asn Glu 40

121 AAC CCT CTG GGA CCT TCT CCC AAG GCC GTG GCG GCG CTG GAG GGA CTG GAC CAC TGG CAC 180
41 Asn Pro Leu Gly Pro Ser Pro Lys Ala Val Ala Ala Leu Glu Gly Leu Asp His Trp His 60

181 CTT TAC CCA GAA GGC TCA AGC TAT GAG CTA CGG CAG GCG CTG GGT AAG AAA CTG GAG ATA 240
61 Leu Tyr Pro Glu Gly Ser Ser Tyr Glu Leu Arg Gln Ala Leu Gly Lys Lys Leu Glu Ile 80

241 GAC CCG GAC AGC ATC ATC GTG GGT TGC GGC TCA AGC GAA GTC ATC CAG ATG CTC TCT TTG 300
81 Asp Pro Asp Ser Ile Ile Val Gly Cys Gly Ser Ser Glu Val Ile Gln Met Leu Ser Leu 100

301 GCC CTG CTG GCG CCC GGC GAC GAG GTG GTC ATC CCT GTG CCT ACC TTT CCC CGC TAT GAG 360
101 Ala Leu Leu Ala Pro Gly Asp Glu Val Val Ile Pro Val Pro Thr Phe Pro Arg Tyr Glu 120

361 CCC CTG GCA CGG CTC ATG GGG GCT AAT CCC GTA AAA GTT CCC TTG AAG GAC TAC CGC ATC 420
121 Pro Leu Ala Arg Leu Met Gly Ala Asn Pro Val Lys Val Pro Leu Lys Asp Tyr Arg Ile 140

421 GAT GTG GAG GCA GTG GCC CGA GCC CTT TCC CCC CGT ACC AAG CTG GTC TAC CTA TGC AAC 480
141 Asp Val Glu Ala Val Ala Arg Ala Leu Ser Pro Arg Thr Lys Leu Val Tyr Leu Cys Asn 160

481 CCC AAC AAC CCC ACC GGG ACC ATC GTC ACC CGG GAG GAG GTG GAG TGG TTC TTG GAA AAG 540
161 Pro Asn Asn Pro Thr Gly Thr Ile Val Thr Arg Glu Glu Val Glu Trp Phe Leu Glu Lys 180

541 GCG GGG GAG GGG GTT CTC ACC GTG CTG GAC GAG GCC TAC TGC GAG TAC GTG ACC AGC CCC 600
181 Ala Gly Glu Gly Val Leu Thr Val Leu Asp Glu Ala Tyr Cys Glu Tyr Val Thr Ser Pro 200

601 GCC TAC CCT GAT GGG CTC GAT TTC CTG CGC CGG GGC TAC AAT GTG GTG GTG CTG CGC ACC 660
201 Ala Tyr Pro Asp Gly Leu Asp Phe Leu Arg Arg Gly Tyr Asn Val Val Val Leu Arg Thr 220

661 TTC TCC AAG ATC TAC GGG CTG GCC GGG CTG CGC ATA GGG TAC GGT GTG GCG GAC AGG GAG 720
221 Phe Ser Lys Ile Tyr Gly Leu Ala Gly Leu Arg Ile Gly Tyr Gly Val Ala Asp Arg Glu 240

721 CTG GTG GCG GAA CTG CAC CGG GTG CGG GAG CCT TTC AAT GTC AGT TCC GCT GCT CAG ATA 780
241 Leu Val Ala Glu Leu His Arg Val Arg Glu Pro Phe Asn Val Ser Ser Ala Ala Gln Ile 260

781 GCC GCC CTG GCC GCC CTG GAA GAC GAA GAG TTC GTG GCG CTT TCG CGC CAG GTC AAC GAA 840
261 Ala Ala Leu Ala Ala Leu Glu Asp Glu Glu Phe Val Ala Leu Ser Arg Gln Val Asn Glu 280

841 GAA GGG AAG GTT TTT CTC TAC CGA GAA CTG GAG AGG CGG GGG ATC GCC TAC GTG CCC ACC 900
281 Glu Gly Lys Val Phe Leu Tyr Arg Glu Leu Glu Arg Arg Gly Ile Ala Tyr Val Pro Thr 300

901 GAG GCC AAC TTC CTA CTC TTC GAT GCC GGT CGG GAC GAG CAG GAA GTA TTT CGC CGG ATG 960
301 Glu Ala Asn Phe Leu Leu Phe Asp Ala Gly Arg Asp Glu Gln Glu Val Phe Arg Arg Met 320

961 CTG CGC CAG GGA GTG ATC ATC CGG GNC GGG GTG GGT TAT CCC ACC CAC TTA AGG GTG ACC 1020
321 Leu Arg Gln Gly Val Ile Ile Arg Xxx Gly Val Gly Tyr Pro Thr His Leu Arg Val Thr 340

1021 ATC GGC ACC TTG GAA CAG AAC CAG CGC TTC CTG GAA GCT TTG GAT AAG GCT CTA GAG CTT 1080
341 Ile Gly Thr Leu Glu Gln Asn Gln Arg Phe Leu Glu Ala Leu Asp Lys Ala Leu Glu Leu 360

1081 AGG GGG GTT TAA 1092
361 Arg Gly Val End 364

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FIGURE 10

Aquifex aspartate aminotransferase

1 ATG AGA AAA GGA CTT GCA AGT AGG GTA AGT CAC CTA AAA CCT TCC CCC ACG CTG ACC ATA 60 39
Met Arg Lys Gly Leu Ala Ser Arg Val Ser His Leu Lys Pro Ser Pro Thr Leu Thr Ile 1/0

61 ACC GCA AAA GCA AAA GAA TTA AGG GCT AAA GGA GTG GAC GTT ATA GGT TTT GGA GCG GGA 120
Thr Ala Lys Ala Lys Glu Leu Arg Ala Lys Gly Val Asp Val Ile Gly Phe Gly Ala Gly

121 GAA CCT GAC TTC GAC ACA CCC GAC TTC ATA AAG GAA GCC TGT ATA AGG GCT TTA AGG GAA 180
Glu Pro Asp Phe Asp Thr Pro Asp Phe Ile Lys Glu Ala Cys Ile Arg Ala Leu Arg Glu

181 GGA AAG ACC AAG TAC GCT CCC TCC GCG GGA ATA CCA GAG CTC AGA GAA GCT ATA GCT GAA 240
Gly Lys Thr Lys Tyr Ala Pro Ser Ala Gly Ile Pro Glu Leu Arg Glu Ala Ile Ala Glu

241 AAA CTA CTG AAA GAA AAC AAA GTT GAG TAC AAA CCT TCA GAG ATA GTC GTT TCC GCA GGA 300
Lys Leu Leu Lys Glu Asn Lys Val Glu Tyr Lys Pro Ser Glu Ile Val Val Ser Ala Gly

301 GCG AAA ATG GTT CTC TTC CTC ATA TTC ATG GCT ATA CTG GAC GAA GGA GAC GAG GTT TTA 360
Ala Lys Met Val Leu Phe Leu Ile Phe Met Ala Ile Leu Asp Glu Gly Asp Glu Val Leu

361 CTA CCT AGC CCT TAC TGG GTA ACT TAC CCC GAA CAG ATA AGG TTC TTC GGA GGG GTT CCC 420
Leu Pro Ser Pro Tyr Trp Val Thr Tyr Pro Glu Gln Ile Arg Phe Phe Gly Gly Val Pro

421 GTT GAG GTT CCT CTA AAG AAA GAG AAA GGA TTT CAA TTA AGT CTG GAA GAT GTG AAA GAA 480
Val Glu Val Pro Leu Lys Lys Glu Lys Gly Phe Gln Leu Ser Leu Glu Asp Val Lys Glu

481 AAG GTT ACG GAG AGA ACA AAA GCT ATA GTC ATA AAC TCT CCG AAC AAC CCC ACT GGT GCT 540
Lys Val Thr Glu Arg Thr Lys Ala Ile Val Ile Asn Ser Pro Asn Asn Pro Thr Gly Ala

541 GTT TAC GAA GAG GAG GAA CTT AAG AAA ATA GCG GAG TTT TGC GTG GAG AGG GGC ATT TTC 600
Val Tyr Glu Glu Glu Glu Leu Lys Lys Ile Ala Glu Phe Cys Val Glu Arg Gly Ile Phe

601 ATA ATT TCC GAT GAG TGC TAT GAG TAC TTC GTT TAC GGT GAT GCA AAA TTT GTT AGC CCT 660
Ile Ile Ser Asp Glu Cys Tyr Glu Tyr Phe Val Tyr Gly Asp Ala Lys Phe Val Ser Pro

661 GCC TCT TTC TCG GAT GAA GTA AAG AAC ATA ACC TTC ACG GTA AAC GCC TTT TCG AAG AGC 720
Ala Ser Phe Ser Asp Glu Val Lys Asn Ile Thr Phe Thr Val Asn Ala Phe Ser Lys Ser

721 TAT TCC ATG ACT GGT TGG CGA ATA GGT TAT GTA GCG TGC CCC GAA GAG TAC GCA AAA GTG 780
Tyr Ser Met Thr Gly Trp Arg Ile Gly Tyr Val Ala Cys Pro Glu Glu Tyr Ala Lys Val

781 ATA GCG AGT CTT AAC AGC CAG AGT GTT TCC AAC GTC ACT ACC TTT GCC CAG TAT GGA GCT 840
Ile Ala Ser Leu Asn Ser Gln Ser Val Ser Asn Val Thr Thr Phe Ala Gln Tyr Gly Ala

841 CTT GAG GCC TTG AAA AAT CCA AAG TCT AAA GAT TTT GTA AAC GAA ATG AGA AAT GCT TTT 900
Leu Glu Ala Leu Lys Asn Pro Lys Ser Lys Asp Phe Val Asn Glu Met Arg Asn Ala Phe

901 GAA AGG AGA AGG GAT ACG GCT GTA GAA GAG CTT TCT AAA ATT CCA GGT ATG GAT GTG GTA 960
Glu Arg Arg Arg Asp Thr Ala Val Glu Glu Leu Ser Lys Ile Pro Gly Met Asp Val Val

961 AAA CCC GAA GGT GCC TTT TAC ATA TTT CCG GAC TTC TCC GCT TAC GCT GAG AAA CTG GGT 1020
Lys Pro Glu Gly Ala Phe Tyr Ile Phe Pro Asp Phe Ser Ala Tyr Ala Glu Lys Leu Gly

1021 GGT GAT GTG AAA CTC TCG GAG TTC CTT CTG GAA AAG GCT AAG GTT GCG GTG GTT CCC GGT 1080
Gly Asp Val Lys Leu Ser Glu Phe Leu Leu Glu Lys Ala Lys Val Ala Val Val Pro Gly

1081 TCG GCC TTC GGA GCT CCC GGA TTT TTG AGG CTT TCT TAC GCC CTT TCC GAG GAA AGA CTC 1140
Ser Ala Phe Gly Ala Pro Gly Phe Leu Arg Leu Ser Tyr Ala Leu Ser Glu Glu Arg Leu

1141 GTT GAG GGT ATA AGG AGA ATA AAG AAA GCC CTT GAA GAG ATC TAA 1185
Val Glu Gly Ile Arg Arg Ile Lys Lys Ala Leu Glu Glu Ile End

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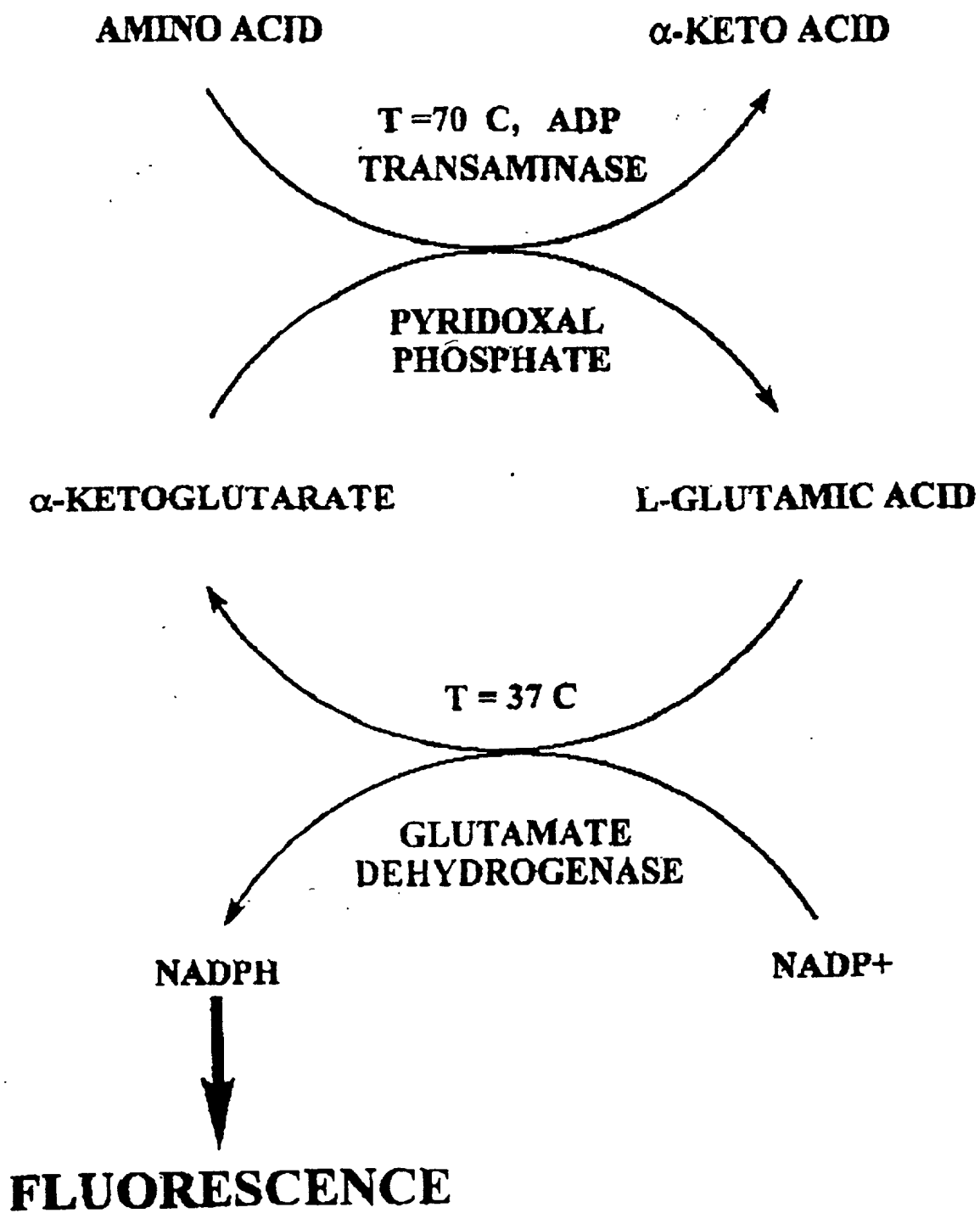


Figure 11

DECLARATION AND POWER OF ATTORNEY

As a below named inventor, I hereby declare that:

My residence, post office address and citizenship are as stated below next to my name.

I believe I am the original, first and sole inventor (if only one name is listed below) or an original, first and joint inventor (if plural names are listed below) of the subject matter which is claimed and for which a patent is sought on the invention entitled:

TRANSAMINASES AND AMINOTRANSFERASES

the specification of which [] is attached hereto or [X] was filed on May 8, 1996 as Application Serial No. 08/656,590 and was amended on _____ (if applicable).

I hereby state that I have reviewed and understand the contents of the above identified specification, including the claims, as amended by any amendment referred to above.

I acknowledge the duty to disclose information which is material to the patentability of this application in accordance with Title 37, Code of Federal Regulations, Section 1.56(a).

I hereby claim foreign priority benefits under Title 35, United States Code, Section 119 of any foreign application(s) for patent or inventor's certificate listed below and have also identified below any foreign application for patent or inventor's certificate having a filing date before that of the application on which priority is claimed. Prior Foreign Application(s):

			Priority Claimed	
(Number)	(Country)	(Day/Month/Year Filed)	Yes	No
			<input type="checkbox"/>	<input type="checkbox"/>

I hereby claim the benefit under Title 35, United States Code, Section 120 of any United States application(s) listed below and, insofar as the subject matter of each of the claims of this application is not disclosed in the prior United States application in the manner provided by the first paragraph of Title 35, United States Code, Section 112, I acknowledge the duty to disclose material information as defined in Title 37, Code of Federal Regulations, Section 1.56(a) which occurred between the filing date of the prior application and the national or PCT international filing date of this application:

<u>08/599,171</u>	<u>February 9, 1996</u>	<u>Pending</u>
(Application Serial No.)	(Filing Date)	(Status - patented, pending, abandoned)
<u> </u>	<u> </u>	<u> </u>
(Application Serial No.)	(Filing Date)	(Status - patented, pending, abandoned)

I hereby appoint the following attorney(s) and/or agent(s) to prosecute this application and to transact all business in the Patent and Trademark Office connected therewith: John N. Bain (Reg. No. 18,651); John G. Gilfillan, III (Reg. No. 22,746); Elliot M. Olstein (Reg. No. 24,025); Raymond J. Lillie (Reg. No. 31,778); Charles J. Herron (Reg. No. 28,019); William Squire (Reg. No. 25,378); and J.G. Mullins (Reg. No. 33,073). Address correspondence and telephone calls to Charles J. Herron c/o Carella, Byrne, Bain, Gilfillan, Cecchi, Stewart & Olstein, 6 Becker Farm Road, Roseland, NJ 07068 - (201) 994-1700.

I hereby declare that all statements made herein of my own knowledge are true and that all statements made on information and belief are believed to be true; and further that these statements were made with the knowledge that willful false statements and the like so made are punishable by fine or imprisonment, or both, under Section 1001 of Title 18 of the United States Code and that such willful false statements may jeopardize the validity of the application or any patent issued thereon.

Full name of inventor: Patrick V. Warren

Inventor's signature: Patrick V. Warren

Date: 7/21/96

Residence: 3507 Sheffield Avenue, Philadelphia, PA 19163

Citizenship: United States

Post Office Address: same

Full name of inventor: Ronald V. Swanson

Inventor's signature: Ronald V. Swanson

Date: 7/23/96

Residence: 309 No. Lemon Street, Apt. A, Media, PA 19063

Citizenship: United States

Post Office Address: same

INSTITUTE DECLARATION AND POWER OF ATTORNEY

As a below named inventor, I hereby declare that:

My residence, post office address and citizenship are as stated below next to my name.

I believe I am the original, first and sole inventor (if only one name is listed below) or an original, first and joint inventor (if plural names are listed below) of the subject matter which is claimed and for which a patent is sought on the invention entitled:

TRANSAMINASES AND AMINOTRANSFERASES

the specification of which [] is attached hereto or [X] was filed on May 8, 1996 as Application Serial No. 08/646,590 and was amended on _____ (if applicable).

I hereby state that I have reviewed and understand the contents of the above identified specification, including the claims, as amended by any amendment referred to above.

I acknowledge the duty to disclose information which is material to the patentability of this application in accordance with Title 37, Code of Federal Regulations, Section 1.56(a).

I hereby claim foreign priority benefits under Title 35, United States Code, Section 119 of any foreign application(s) for patent or inventor's certificate listed below and have also identified below any foreign application for patent or inventor's certificate having a filing date before that of the application on which priority is claimed. Prior Foreign Application(s):

(Number)	(Country)	(Day/Month/Year Filed)	Priority Claimed Yes No <input type="checkbox"/> <input type="checkbox"/>
08/599,171	February 9, 1996	Pending	
(Application Serial No.)	(Filing Date)	(Status - patented, pending, abandoned)	
(Application Serial No.)	(Filing Date)	(Status - patented, pending, abandoned)	

I hereby appoint the following attorney(s) and/or agent(s) to prosecute this application and to transact all business in the Patent and Trademark Office connected therewith: John N. Bain (Reg. No. 18,651); John G. Gilfillan, III (Reg. No. 22,746); Elliot M. Olstein (Reg. No. 24,025); Raymond J. Lillie (Reg. No. 31,778); Charles J. Herron (Reg. No. 28,019); William Squire (Reg. No. 25,378); and J.G. Mullins (Reg. No. 33,073). Address correspondence and telephone calls to Charles J. Herron c/o Carella, Byrne, Bain, Gilfillan, Cecchi, Stewart & Olstein, 6 Becker Farm Road, Roseland, NJ 07068 - (201) 994-1700.

I hereby declare that all statements made herein of my own knowledge are true and that all statements made on information and belief are believed to be true; and further that these statements were made with the knowledge that willful false statements and the like so made are punishable by fine or imprisonment, or both, under Section 1001 of Title 18 of the United States Code and that such willful false statements may jeopardize the validity of the application or any patent issued thereon.

Full name of inventor: Patrick V. Warren

Inventor's signature: Patrick V. Warren

Date: 12/26/96

Residence: 3507 Sheffield Avenue, Philadelphia, PA 19163

Citizenship: United States

Post Office Address: same

Full name of inventor: Ronald V. Swanson

Inventor's signature: Ronald V. Swanson

Date: 12/23/96

Residence: 309 No. Lemon Street, Apt. A, Media, PA 19063

Citizenship: United States

Post Office Address: same

IN THE UNITED STATES PATENT AND TRADEMARK OFFICE

Applicants: Warren, P.V. *et al.* Art Unit:
Serial No.: 08/646,590 Examiner:
Filed: 5/8/96
Title: TRANSAMINASES AND AMINOTRANSFERASES

Assistant Commissioner for Patents
Washington, DC 20231

REVOCATION AND NEW POWER OF ATTORNEY

Under 37 CFR §3.73(b) RECOMBINANT BIOCATALYSIS, INC., a Delaware corporation, certifies that it is the assignee of 100% of the right, title and interest in the patent application identified above by virtue of an assignment from the inventors of the patent application identified above. A copy of the Assignment, executed on July 23, 1996, is attached hereto.

The undersigned has reviewed all the documents in the chain of title of the patent application identified above and, to the best of undersigned's knowledge and belief, title is in the assignee identified above.

The undersigned, whose title is supplied below, is empowered to act on behalf of the assignee.

Date of Deposit 6-13-97
I hereby certify under 37 CFR 1.8(a) that this correspondence is being deposited with the United States Postal Service as first class mail with sufficient postage on the date indicated above and is addressed to the Assistant Commissioner for Patents, Washington, D.C. 20231.

Stephanie Sharrett
Stephanie Sharrett

The undersigned, acting on behalf of the assignee, hereby revokes all powers of attorney previously granted in the application and appoints: John R. Wetherell, Jr., Ph.D. (Reg. No. 31,678); Lisa A. Haile (Reg. No. 38,347); Stacy L. Taylor (Reg. No. 34,842); John Land (Reg. No. 29,554), and June M. Learn (Reg. No. 31,238), of the firm of FISH & RICHARDSON P.C., as its attorneys with full power of substitution and revocation, to prosecute the application and to transact all business in the United States Patent and Trademark Office connected therewith.

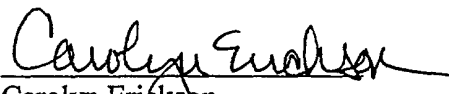
Please direct all telephone calls to John R. Wetherell at (619) 678-5070 and all correspondence relative to said application to the following address:

John R. Wetherell, Jr., Ph.D.
FISH & RICHARDSON, P.C.
4225 Executive Square, Suite 1400
La Jolla, California 92037
(619) 678-5099 (Facsimile)

I hereby declare that all statements made herein of my own knowledge are true and that all statements made on information and belief are believed to be true; and further that these statements were made with the knowledge that willful false statements and the like so made are punishable by fine or imprisonment, or both, under Section 1001 of Title 18 of the United States Code and that such willful false statements may jeopardize the validity of the application or any patents issued thereon.

Respectfully submitted,

Date: 5-16-97


Carolyn Erickson
Manager, Business Development and
Regulatory Affairs
RECOMBINANT BIOCATALYSIS, INC.